One Panel.

Three Major Infectious Causes.

Real-Time PCR testing for HSV-1/HSV-2, H. ducreyi, and T. pallidum from a single specimen.

Why Order the MDL Genital Ulcer Disease Panel?

Supports Differential Diagnosis

Helps distinguish herpes, syphilis, and chancroid when clinical presentation alone is inconclusive.

Identifies HSV Subtype

Differentiates HSV-1 from HSV-2 to support counseling and clinical management.

Sensitive Molecular Detection

Real-Time PCR provides direct detection of pathogen DNA.

Simplifies Specimen Collection

Utilizes MDL's ***OneSwab***® or ThinPrep® specimen collection platforms, supporting a streamlined workflow for your practice.

Test No. 115 | Genital Ulcer Disease Panel

HSV-1 & HSV-2 • *Haemophilus ducreyi* • *Treponema pallidum* (syphilis)

Single specimen collection. Single panel order. Comprehensive results.

When clinical presentation isn't enough.

Consider MDL Test No. 115 when genital ulcer disease is suspected and the infectious cause is unclear.

A MEMBER OF GENESIS BIOTECHNOLOGY GROUP

Medical Diagnostic Laboratories
www.mdlab.com • 877.269.0090

A DIVISION OF GENESIS CLINICAL DIAGNOSTICS

IH0327 Rel.: 8.2026

Medical Diagnostic Laboratories
2439 Kuser Road
Hamilton, NJ 08690
Toll Free: 877-269-0090
Fax: 609-570-1050
www.mdlab.com

PATIENT

MDL # 13364148

FINAL

DOE, JANE
555 MAIN ROAD
ANYTOWN, NJ 12345-6789
DOB: 11/30/1998 (Age 25)
Gender: Female
Ethnicity: Not provided
Patient ID: 82100
Home #: 123-456-7890

SPECIMEN

Type	Source	Collected	Received	Reported
Swab	Vag/Anal	04/15/2024	04/16/2024	04/19/2024

CLIENT **NPI:** 0987654321

DOE FAMILY PRACTICE
JOHN DOE, MD
1234 FIRST AVENUE
ANYTOWN, NJ 12345-6789
Tel: (555) 555-1234
Fax: 555-555-1235

Pathogens Detected

126 Herpes subtype (HSV-2) *	POSITIVE	110 *Treponema pallidum* (syphilis) *	POSITIVE

Pathogens Not Detected

122 *Haemophilus ducreyi* *	126 Herpes subtype (HSV-1) *

*This test was developed and its performance characteristics determined by the laboratory. It has not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary.

A positive result is provided for bacteria, virus, parasites, and/or fungal species when PCR amplification (real-time PCR), sequence information (Pyrosequencing), and/or sequencing analysis occurs above cut-off levels established by the laboratory. Pertinent reference intervals for the tests reported above are available from the laboratory upon request.

Medical Director, Jing-Jing Yang, M.D.

MDL#: **13364148** 05/20/2024

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