CIRRUSDX

FINAL

Accession #: S2025-000317
Name: Doe, Jane
DOB: 1/6/1999
MRN/SSN:
Sex: Female

Laboratory Information	Sample Information	Practice Information
CirrusDx, Inc 77 Upper Rock Circle, 4th Floor Rockville, MD 20850 CLIA# 21D2130541 Dr. Todd Myers, Lab Director 240-813-8801 or reports@cirrusdx.com	**Technician:** CWS **Date Collected:** 8/6/2025 **Date Received:** 8/7/2025 **Date Reported:** 8/7/2025 **Reporting Method:**	**Name:** Apple Urology **Provider:** Alexandra Mary, Elizabeth **Address:** 123 Apple Way Farmville MD 12345 **Phone:** 987-654-3210 **Fax:** (301) 621-4254 **Email:** results@appleurology.com

Testing Information

Test Name:	Test Method:	Specimen Type:	Specimen Site:
Sexually Transmitted Infection Panel (STIP™)	Real-Time PCR (Molecular)	Swab	Rectal

Pathogen	Result
Chlamydia trachomatis	**Detected**
Neisseria gonorrhoeae	**Not Detected**
Trichomonas vaginalis	**Not Detected**
Ureaplasma spp.	**Not Detected**
Mycoplasma hominis	**Not Detected**
Mycoplasma genitalium	**Not Detected**

General Comments

For treatment guidelines refer to the 'CDC Sexually Transmitted Infections Treatment Guidelines, 2021' document.

Notes:

Questions including Clinical Consultation on Results: 240-813-8801 or reports@cirrusdx.com

Organisms Tested (Reference Range): *Chlamydia trachomatis* (Not Detected), *Neisseria gonorrhoeae* (Not Detected), *Mycoplasma genitalium* (Not Detected), *Mycoplasma hominis* (Not Detected), *Trichomonas vaginalis* (Not Detected), *Ureaplasma spp.* (Not Detected)

It is the physician's responsibility to interpret the results provided and determine the appropriate (if any) treatment options.

The assays were developed and their performance characteristics were determined by CirrusDx. They have not been cleared or approved by the US Food and Drug Administration. The FDA does not require these tests to go through premarket FDA review. This test is used for clinical purposes. It should not be regarded as investigational or for research. This laboratory is certified under the Clinical Laboratory Improvement Amendments (CLIA) as qualified to perform high complexity clinical laboratory testing.

This report has been reviewed and approved by: Todd E Myers

Dr. Todd Myers, Laboratory Director.

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