



PATIENT INFORMATION

Name: Patient Sample
 DOB: 01/01/1975
 Sex at Birth: F
 Address: 123 Resident Street
 City, State, Zip
 Phone: 123.456.7890
 Race:
 Ethnicity:



SPECIMEN INFORMATION

Lab Accession Number: 13044771
 Menu: Vaginitis
 Specimen Type: Vaginal
 Date Collected: 03/15/2026
 Date Received by Lab: 03/16/2026
 Date Reported: 03/16/2026
 Cross Reference #:



FACILITY INFORMATION

Facility Name: Women's OBGYN
 Provider Name: Doctor Test, MD
 NPI: 10000000
 Address: 123 Main Street
 City, State, Zip
 Phone: 555.555.5555

TEST RESULTS

ORGANISMS	TARGETS	DETECTION STATUS	MICROBIAL LOAD
	Bacterial Vaginosis	Positive	

CLINICAL NOTES

- **Bacterial Vaginosis** is reported as a qualitative result from algorithmic based detection of the following organisms: *Lactobacillus crispatus*, *Lactobacillus jensenii*, *Gardnerella vaginalis*, *Atopobium vaginae*, Bacterial Vaginosis Associated Bacteria-2 (BVAB-2), and *Megasphaera-1*.

CONSIDERATIONS FOR THERAPY

The information below lists agents that may be considered for therapy based on detected organisms and/or resistance genes. This information does not constitute medical advice. Clinical management and treatment decisions remain the sole responsibility of the ordering healthcare provider.



References and guidelines used to develop this table.

(P) Primary therapy consideration (A) Alternative therapy consideration (N/A) Not Applicable

CLASS OF ANTIMICROBIAL AGENTS	ANTIBIOTIC (ADMINISTRATION)	Bacterial Vaginosis
Lincosamide	Clindamycin (po/VAG)	A
	Metronidazole (po/VAG)	P
Nitroimidazole	Tinidazole	A
	Secnidazole	A

TARGETS TESTED

Targets detected are bolded for reference

BACTERIAL


Bacterial Vaginosis
 Chlamydia trachomatis
 Neisseria gonorrhoeae

PROTOZOAL

Trichomonas vaginalis

FUNGAL

Candida albicans, parapsilosis, tropicalis
 Candida glabrata (Nakaseomyces glabratus)
 Candida krusei (Pichia kudriavzevii)

Analyzed by: Carrie Wilks, PhD 

Date: 11/04/2025 13:16

Have questions about this report?

Call our Clinical Expert Line at **866-287-3218**.



Real-Time polymerase chain reaction (TaqMan qPCR) was utilized for detection for all tested organisms and resistance genes. Initiation of antimicrobial therapy prior to testing may affect results and can lead to the detection of non-living microorganisms. Detection of microbes must be correlated with current/recent antibiotic usage and patient signs and symptoms. Microbial sensitivity testing is not performed at this lab. Crt to CFU/mL equivalent thresholds were established based on studies using known CFU/mL urine specimens performed at HealthTrackRx in Denton, TX. Testing performed by HealthTrackRx at LabPort (2425 Universal Way Ste 6, Louisville, KY 40219; CLIA# 18D2319534; Lab Director Jessica Moye, PhD, HCLD(ABB)). Data analysis and report release for all tested samples is referred out to HealthTrackRx Indiana, Inc (1500 I35W, Denton, TX 76207; CLIA# 45D2009077) and performed by their appropriately credentialed staff according to CAP and Texas State Requirements. This test was developed, and its performance characteristics determined by HealthTrackRx. This test was developed, and its performance characteristics determined by HealthTrackRx. It has not been cleared or approved by the FDA. However, such approval/clearance is not required, as the laboratory is regulated and qualified under CLIA to perform high-complexity testing. This test is used for clinical purposes and should not be regarded as investigational or for research.

*Approximate copies of target nucleic acid per μL (Low: $<2,500$ copies/ μL ; Moderate: 2,500-50,000 copies/ μL ; High: $>50,000$ copies/ μL)

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