

Patient: Test, Patient MRN: 4 Accession: 18919

Patient #: 6818 Birth: 4/26/1950

 Doctor:
 Test, Doctor
 Age:
 73 years
 Collection Date:
 10/4/2023 2:29 PM
 TB

 Home Phone:
 1231231234
 Gender:
 Male
 Received Date:
 10/4/2023 2:29 PM
 TB

Specimen Source: Urine

Test Name	Result	Units	Flag	Reference Range/Cutoff
Vaginitis				Run by TB on 10/4/2023 2:32:13 PM
Atopobium vaginae	Not Detected			
Bacteroides fragilis	Detected		Abnormal	
BVAB-2	Not Detected			
Candida albicans	Not Detected			
Candida dubliniensis	Not Detected			
Candida glabrata	Not Detected			
Candida krusei	Detected		Abnormal	
Candida lusitaniae	Not Detected			
Candida parapsilosis	Detected		Abnormal	
Candida tropicalis	Not Detected			
Chlamydia trachomatis	Not Detected			
Enterococcus spp.	Detected		Abnormal	
Escherichia coli	Not Detected			
Gardnerella vaginalis	Not Detected			
Haemophilus ducreyi	Detected		Abnormal	
HHV-1 & 2 (Human Herpes Virus)	Detected		Abnormal	
Lactobacillus crispatcus	Not Detected			
Lactobacillus gasseri	Not Detected			
Lactobacillus iners	Not Detected			
Lactobacillus jensenii	Detected		Abnormal	
Megasphaera Type 1	Not Detected			
Megasphaera Type 2	Not Detected			
Mobiluncus curtisii	Detected		Abnormal	
Mobiluncus mulieris	Detected		Abnormal	
Mycoplasma genitalium	Detected		Abnormal	
Mycoplasma hominis	Not Detected			
Neisseria gonorrhoeae	Detected		Abnormal	
Prevotella bivia	Not Detected			
Staphylococcus aureus	Not Detected			
Streptococcus agalactiae (Grp B)	Not Detected			
Treponema pallidum	Not Detected			
Trichomonas vaginalis	Detected		Abnormal	
Ureaplasma urealyticum	Not Detected			

Originally Reported On: 10/4/2023 2:34 PM

Printed: 1/10/2024 1:27 PM

(UTC-06:00) Central Time (US & Canada)

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STAT[S] Corrected [C] Amended [A]



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Notes:

Disclaimer: Testing was performed at OmniHealth Diagnostics, LLC (CLIA# 45D2089485) Laboratory Director Akhtar Afshan Ali. This is a Laboratory Developed Test, and its performance characteristics are determined by OmniHealth Diagnostics, LLC. This laboratory is regulated under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) and is qualified to perform high-complexity clinical testing. This test has not been cleared nor approved by the U.S. Food and Drug Administration (FDA).

Method: Nucleic acid from patient specimens was amplified by RT-PCR using reagents and control materials validated by OmniHealth Diagnostics, LLC for the detection of selected microorganisms and Antibiotic Resistance Markers.

Test Result Interpretation:

Detected: Indicates the presence of the target microorganism(s).

Inconclusive: A small amount of nucleic acid was identified but inadequate to confidently make a result call. This result can occur from inadequate sample collection, new or very early-stage infection, or for patients close to recovery. Recollection and Retesting recommended.

Invalid: Unable to confirm the presence or absence of the targeted microorganism due to specimen quality. Recollection and Retesting recommended.

Not Detected: A negative detection does not imply the absence of microorganisms other than those listed or does not exclude the possibility that the target sequence is present below the limit of detection.

It is recommended that the test results be interpreted in conjunction with all other medical information relevant to the patient's clinical condition

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