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|-------------|---------------|---------|-----------|------------------|-----------------------|
| Patient: | Test, Patient | MRN: | 4 | Accession: | 18883 |
| Patient #: | 6818 | Birth: | 4/26/1950 | | |
| Doctor: | Test, Doctor | Age: | 73 years | Collection Date: | 10/4/2023 10:42 AM TB |
| Home Phone: | 1231231234 | Gender: | Male | Received Date: | 10/4/2023 10:42 AM TB |
| | | | | Specimen Source: | Urine |

| Test Name | Result | Units | Flag | Reference Range/Cutoff |
|--------------------------------|----------|-------|-----------------|--|
| STI | | | | <i>Run by TB on 10/4/2023 1:55:57 PM</i> |
| HHV-1 & 2 (Human Herpes Virus) | Detected | | Abnormal | |
| Atopobium vaginae | Detected | | Abnormal | |
| Chlamydia trachomatis | Detected | | Abnormal | |
| Gardnerella vaginalis | Detected | | Abnormal | |
| Haemophilus ducreyi | Detected | | Abnormal | |
| Neisseria gonorrhoeae | Detected | | Abnormal | |
| Treponema pallidum | Detected | | Abnormal | |
| Trichomonas vaginalis | Detected | | Abnormal | |

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

Accession: 18883 Patient: #: 6818

Originally 1 Rep/2023 D28 PM 4/2023 1:59 PM

Lab Results for: Test, Patient

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

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

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