
Lab Dept: Urine/Stool

Test Name: BLASTOMYCES ANTIGEN, URINE

General Information

Lab Order Codes: BAG

Synonyms: N/A

CPT Codes: 87449 – Infectious agent antigen detection by EIA technique

Test Includes: Urine Blastomyces Antigen level reported in ng/mL.

Logistics

Test Indications: Diagnosis of infection with Blastomyces dermatidis. Monitor antigen levels following initiation of antifungal treatment.

Lab Testing Sections: Urine/Stool - Sendouts

Referred to: Mayo Clinic Laboratories (MML Test: UBLAS)

Phone Numbers: MIN Lab: 612-813-6280

STP Lab: 651-220-6550

Test Availability: Daily, 24 hours, test performed Monday - Saturday

Turnaround Time: 1 -2 days

Special Instructions: Use sterile collection technique.

Specimen

Specimen Type: Urine, random

Container: Sterile leak-proof container

Draw Volume: 4 mL (Minimum: 2.5 mL) urine

Processed Volume: Same as Draw Volume

Collection: Routine urine collection, or other appropriate collection technique. Maintain sterility during collection and forward promptly to the laboratory.

Special Processing: Lab Staff: Mix random urine sample well. Remove aliquot into a plastic scew-capped tube.. Store and ship at refrigerated temperatures. Forward promptly.

Patient Preparation: None

Sample Rejection: Mislabeled or unlabeled specimens

Interpretive

| Reference Range: | ng/mL | Result Interpretation |
|-------------------------|------------------|------------------------------|
| | 0.0 mg/dL | None Detected |
| | <1.3 ng/mL | Detected |
| | 1.3 – 20.0 ng/mL | Detected |
| | >20.0 ng/mL | Detected |

Critical Values: N/A

Limitations:

- Cross-reactivity with other fungal infections, including *Histoplasma capsulatum*, may occur. Positive results should be correlated with other clinical and laboratory findings (eg, culture, serology)
- Low-level positive antigen levels may persist following resolution of infection and completion of appropriate treatment regimen
- Turbid urine specimens, containing excess protein, cells or particulate matter can inhibit the function of the test
- Sensitivity of this assay to detect antigen from species other than *Blastomyces dermatidis* is unknown

Methodology: EIA (Enzyme Immunoassay)

References: [Mayo Clinic Laboratories](#) May 2021

Updates:

7/2/2009: Added other specimen types (BALF, other sterile body fluids)
2/8/2011: Units change, previously reported as EIA Units. Reference range update.
5/15/2012: Note reference range interpretation update.
1/13/2015: Added plasma as a specimen type.
3/9/2020: Orderable test is specific to Urine, other specimen types removed.
5/4/2021: No longer forwarded to Mira Vista, Mayo performing internally.