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AFLATOXIN GROUPS (B1, B2, B4, B4) REPORT FORM

Patient:

Patient Date of Birth:

Accession No:

Date of Service:

Patient ID No:

Date of Report:

Collected:

Ordering Physician: Dr. Anna Davis-DirectLabs

Procedure:

TYPE: Aflatoxin (Procedure by ELISA).

Test Results:

Code	Test	Specimen	Value	Result	Negative if less than	Equivocal if between	Positive if greater or equal
E8502	Aflatoxin	Urine	0 ppb	Negative	0.8 ppb	0.8-1.0 ppb	1.0 ppb

Tests such as this should be used only in conjunction with other medically established diagnostic elements (e.g., symptoms, history, clinical impressions, results from other tests, etc). Physicians should use all the information available to them to diagnose and determine appropriate treatment for their patients.

Director's Initials: JSS

Disclaimer: This test was developed and its performance characteristics determined by RealTime Lab. It has not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. 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 of 1988 (CLIA-88) as qualified to perform high complexity clinical laboratory testing.