Patient NAME	:		
DOB/Age/Gender	:	Report STATUS	:
Patient ID / UHID	:	Barcode NO	:
Referred BY	:	Sample Type	:
Sample Collected	:	Report Date	:

Galactomannan (Aspergillus Antigen)

Test Description	Value(s)	Unit(s)	Reference Range
GALACTOMANNAN (ASPERGILLUS ANTIGEN) *EIA, Serum/Bal Fluid*	0.78	Index	Negative: < 0.9 Equivocal: 0.9-1.1 Positive: > 1.1

Interpretation:
1. In case of equivocal result , please send a repeat sample after 2-4 weeks. If result remains equivocal , the same should be considered negative.
2. Galactomannan antigen test does not rule out a diagnosis of invasive aspergillosis ; clinical correlation is recommended.

Comments:-
Detection of Aspergillus genus-specific antigens is important for diagnosis of life threatening invasive pulmonary aspergillosis. This form of aspergillosis is usually observed in patients with primary or secondary immunodeficiency. Galactomannan is an oligosaccharide determinant specific for Aspergillus genus;its detection allows to distinguish aspergillosis from other mycoses. Galactomannan antigen of Aspergillus spp. Fungi may be used for diagnosis of invasive aspergillosis in combination with microbiological , histological and Xray data.

*** End Of Report ***

Terms and Conditions of Reporting

1. The presented findings in the Reports are intended solely for informational and interpretational purposes by the referring physician or other qualified medical professionals possessing a comprehensive understanding of reporting units, reference ranges, and technological limitations. The laboratory shall not be held liable for any interpretation or misinterpretation of the results, nor for any consequential or incidental damages arising from such interpretation.

2. It is to be presumed that the tests performed pertain to the specimen/sample attributed to the Customer's name or identification. It is presumed that the verification particulars have been cleared out by the customer or his/her representation at the point of generation of said specimen / sample. It is hereby clarified that the reports furnished are restricted solely to the given specimen only.

3. It is to be noted that variations in results may occur between different laboratories and over time, even for the same parameter for the same Customer. The assays are performed and conducted in accordance with standard procedures, and the reported outcomes are contingent on the specific individual assay methods and equipment(s) used, as well as the quality of the received specimen.

4. This report shall not be deemed valid or admissible for any medico-legal purposes.

5. The Customers assume full responsibility for apprising the Company of any factors that may impact the test finding. These factors, among others, includes dietary intake, alcohol, or medication / drug(s) consumption, or fasting. This list of factors is only representative and not exhaustive.