

Patient Name :
 DOB/Age/Gender : Bill Date :
 Patient ID / UHID : Sample Collected :
 Referred By : Sample Received :
 Sample Type : Report Date :
 Barcode No : Report Status :

Test Description	Value(s)	Unit(s)	Reference Range
------------------	----------	---------	-----------------

BIOCHEMISTRY REPORT

Progesterone

Progesterone 7.64 ng/ml
 Method : ECLIA

Interpretation:

Healthy men	< 0.050 - 0.149
Healthy postmenopausal women	< 0.050 - 0.126
Healthy pregnant women	
1st trimester	11.0 - 44.3
2nd trimester	25.4 - 83.4
3rd trimester	58.7 - 214.0
Healthy women Cycle Phase	
Follicular	< 0.050 - 0.193
Ovulation	0.055 - 4.14
Luteal	4.11 - 14.5
Healthy women Cycle Sub Phase	
Early follicular	< 0.050 - 0.323
Intermediate follicula	< 0.050 - 0.22
Late follicular	< 0.050 - 0.216
Ovulation	< 0.050 - 2.35
Early luteal	2.36 - 15.1
Intermediate luteal	4.76 - 20.9
Late luteal	0.537 - 13.5




Dr. Rosy Khandelia
MD Pathology
Consultant Pathologist

Booking Centre :- HEALTH CARE DIAGNOSTIC CENTRE (GOLAGHAT), Sajjad Ahmed

Processing Lab :- Redcliffe Lifetech Pvt. Ltd., Second Floor, Dag No.238, 243, Patta no.723,1166, Mauza, Beltola - Khanapara Rd, Odalbakra, Guwahati, Assam 781034

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.

DISCLAIMER

This is a sample report provided for demonstration purposes only and does not represent an actual patient report. Test results, reference ranges, methodologies, instrumentation, and report formats may vary depending on the laboratory performing the test. The format and representation shown are indicative of reports generated by the National Reference Laboratory of Redcliffe Labs, Noida. This sample report should not be used for medical interpretation, diagnosis, or treatment decisions.