LABORATORY REPORT

Redcliffe labs

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

Test Description	Value(s)	Unit(s)	Reference Range
Indirect Coombs Test (ICT)			
Coombs Test, Indirect *Column agglutination technology*	Negative	-	Negative

Interpretation:

Result	
Negative	No Agglutination Seen
Positive	Agglutination Seen

Comment:

1. ICT is used to detect red cells sensitized with IgG antibodies, IgG autoantibodies or complemant components.
2. Negative result means that the mother has not developed antibodies against the fetuss blood. A negative Coombs test indicates that the fetus is not presently in danger from problems relating to Rh incompatibility.
3. Positive result means that the mother has developed antibodies to the fetal red blood cells and is sensitized. However, a positive Coombs test only indicates that an Rh-positive fetus has a possibility of being harmed. A positive test cannot indicate the amount of fetal harm that has occurred or is likely to occur.

*** 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.