LABORATORY REPORT

Redcliffe labs

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

Test Description	Value(s)	Unit(s)	Reference Range
Reticulocyte Count			
Reticulocyte Count *Supravital stain- Brilliant Cresyl blue.*	2.1	%	0.5-2.5

Interpretation:

Reticulocyte count is increased in:

1. Hemolytic anemia.
2. Nutritional anemia after treatment.
3. Acute and chronic blood loss
4. Pregnancy.
5. Bone marrow regeneration.

Reticulocyte count is decreased in:

1. Untreated nutritional anemia.
2. Aplastic anemia
3. Exposure to radiation or radiation therapy.
4. Chronic infection.
5. Chemotherapy.

*** 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.