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

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

Hepatitis C Antibody (HCV), Rapid Card

HEPATITIS C ANTIBODY (Anti-HCV) *Qualitative immunoassay,rapid card*	NON REACTIVE		NON REACTIVE

NAME CHANGE AS PR CLIENT REQUEST

Interpretation:

RESULTS	REMARKS
Reactive	Reactive test result indicates presence of Hepatitis C virus infection
Non Reactive	Non Reactive test result indicates absence of Hepatitis C virus infection

NOTE

1.The **4TH Generation** HCV TRI-DOT detects anti-HCV in human serum or plasma and is **only a screening test**. All reactive samples should be confirmed by supplemental assays like RIBA .Therefore for a definitive diagnosis, the patient's clinical history ,symptomatalogy as well as serological data, should be considered. The results should be reported only after complying with above procedure.
2.A non reactive-results does not exclude the possibility of exposure to or infection with HCV.
3.Repeated false results may occur due to non-specefic binding of the sample to the membrane.
4.The presence of anti-HCV does not imply a HepatitisC infection but may be indicative of recent and /or past infection By HCV.
5.Patients with auto-immune liver diseases may show falsely reactive results.
6. False positive results may be observed in patients receiving mouse monoclonal antibodies, on heparin therapy, on biotin supplements for diagnosis or therapy or presence of heterophilic antibodies in serum.
5. False negative reaction may be due to processing of sample collected early in the course of disease, Prozone phenomenon, Immunosuppression & Immuno-incompetence.

Uses ·
To diagnose suspected HCV infection in risk group.
Prenatal Screening of pregnant women and pre surgical/interventional procedures work up.

Note: This is a sample report for illustrative purpose only. Actual report may vary

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.