

Reference No.	: - 2107003804	Age/Gender	: 57 Yrs/Female	
Pt's Name	: Mrs. SARABJEET		NOD 260	
Referred By	: NA		NOD-300	
Sample Collection Date/Time	: 03-Jul-2021	Date	:03-Jul-2021	
Sample Receiving Date/Time	: 04-Jul-2021 05:27AM	Approved Date	:05-Jul-2021 06:13PM	
Sample From	: 360 DIAGNOSTIC & HEALTH SERVICES PVT LTD	Report Print Time	:07-Jul-2021 11:59AM	

Molecular Biology			
Test Description	Observed Value	Biological Reference Interval	
	Hepatitis C RNA Detection by PC	R*	
Hepatitis C RNA Detection by PCR* Real-time Polymerase Chain Reaction	Undetectable or < 10.0	Undetectable or < 10.0 IU/ml	
Method :			
Real Time Polymerase chain Reaction (RT-PCR)			
Hepatitis C is an infectious disease caused by Hepatitis C	C virus (HCV), which can lead to inflammation a	nd significant damage in the liver. Although it predominantly	
infects the cells of the liver, it can also affect other parts	of the body. During the acute phase following th	e initial infection of HCV, it is generally asymptomatic and	
clinically undetectable. About 85 % of the acute infecti	ons become chronic and the remaining naturally	get cured. In rare cases, acute hepatitis is accompanied by	
jaundice, malaise, weakness and anorexia. It is estimated	d that 74 to 86 % of individuals with the acute info	ection develop persistent viremia, which subsequently leads	
to chronic infection and possibly to cirrhosis or hepato	cellular carcinoma. The conventional diagnostic	methods include serological testing and liver biopsy. Since	
HCV cannot be cultured in the clinical laboratory, a sense	sitive molecular testing is needed to confirm the p	presence of the virus such as quantitative real-time PCR.	
Interpretation :			
Sensitivity : 20 IU/ml			
Sensitivity & Dynamic range : 10^1 X 10^10			
A "DETECTED" result will be reported with quantificat	ion in IU/ml. It indicates the degree of active HCV	✓ viral replication in the patient.	
A "LESS THAN DETECTABLE LIMIT" result indic	cates that either absence of HCV RNA in patien	nt~s specimen or HCV RNA level is below the lower limit	
quantification of this assay.			
A "Inconclusive Result" indicates that inhibitory substan	ces may be present in the specimen and collection	n and testing of a new specimen is recommended.	
Conversion Factor : Result (copies/ml) = Result (IU/m	nl) x 3.00		

Methodology details :

* HCV RNA is extracted from plasma by US FDA approved Automatic Extraction machine based on magnetic bead technology.

* Purified RNA is then Amplified and quantified using CE- IVD approved Real time PCR.

* Extraction and Amplification controls (IC) are incorporated in each run to ensure more accurate and precise detection of RNA.

Laboratory is NABL Accredited

*** End Of Report ***



