



Name : Mr. SAMEER AHMAD ELSAMAD Ref No. : 44987

DOB 26/04/1980 Sample No. 2412511259 44 Y / Male Age / Gender **Collected** 09/12/2024 22:09 Referred by DR.ENOMEN Registered 10/12/2024 14:04 CITICARE MEDICAL CENTER Centre Reported 10/12/2024 15:21

BIOCHEMISTRY				
Test	Result Flag	g Unit	Reference Range	Methodology
LIVER FUNCTION TEST				
ALT / SGPT	29	U/L	< or = 50 Please note change. Source: Roche IFU.	IFCC
AST / SGOT	27	U/L	< or = 50 Please note change. Source: Roche IFU.	IFCC
ALP (ALKALINE PHOSPHATASE)	91	U/L	40 - 129 Please note change. Source: Roche IFU.	Colorimetric assay
GGT (GAMMA GLUTAMYL TRANSFERASE)	17	U/L	10 - 71 Please note change. Source: Roche IFU.	Enzymatic colorimetric assay
BILIRUBIN (TOTAL)	0.3	mg/dL	0 - 1.2 Please note change. Source: Roche IFU.	Diazotization
BILIRUBIN (DIRECT)	0.2	mg/dL	0 - 0.2 Please note change. Source: Roche IFU.	Diazotization
INDIRECT BILIRUBIN	0.10	mg/dL	< or = 0.9	Calculated
TOTAL PROTEIN	6.6	g/dL	6.4 - 8.3	Biuret reaction
ALBUMIN (SERUM)	4.3	g/dL	3.5 - 5.2 Please note change. Source: Roche IFU.	Colorimetric assay (Bromocresol Green)
GLOBULIN	2.3	g/dL	2.0 - 3.5	Calculation
A/G RATIO	1.9		0.8 - 2.0	Calculation

Gone V. Shah

Dr. Adley Mark Fernandes Dr. Vyoma V Shah
M.D (Pathology) M.D (Pathology)
Pathologist Clinical Pathologist

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Greeshma P Sidharthan

Test result pertains only to the sample tested and to be interpreted in the light of clinical history. These tests are accredited under ISO 15189:2012 unless specified by (^). Test marked with # is performed in an accredited referral laboratory.





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Mr. SAMEER AHMAD ELSAMAD Name

DR.ENOMEN

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CITICARE MEDICAL CENTER Centre

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IMMUNOLOGY

Unit Test Result Flag **Reference Range** Methodology

HEPATITIS B SURFACE ANTIGEN (HBSAG) COI **ECLIA** 3330 Non-Reactive: < 0.9

> Borderline: =/>0.9 - <1.0 Reactive: =/>1.0

Note changes in method and

reference range. Source: Roche IFU.

Comments: Please correlate clinically

INTERPRETATION NOTES:

A positive HBsAg test result means that the patient is infected with acute or chronic hepatitis B virus or chronic HBV carrier state.

A negative result implies the patient is not infected with hepatitis B.

HEPATITIS C ANTIBODIES 0.05 COI **ECLIA** Non-Reactive: < 0.9

Borderline: =/>0.9 - <1.0

Reactive: =/>1.0

Source: Roche IFU.

INTERPRETATION NOTES:

A non-reactive screening test result does not exclude the possibility of exposure to or infection with HCV. Non-reactive screening results in individuals with prior exposure to HCV may be due to low antibody levels that are below the limit of detection of this assay or lack of reactivity to the HCV antigens used in this assay. Patients with acute or recent HCV infections (< 3 months from time of exposure) may have false-negative HCV antibody results due to the time needed for seroconversion (average of 8 - 9 weeks). Testing for HCV RNA and or RIBA is recommended.

A repeatedly reactive screening result is consistent with current HCV infection, or past HCV infection that has resolved, or biologic false positivity for HCV antibody. Testing for HCV RNA and or RIBA is recommended

HIV I & II ANTIBODY AND P24 ANTIGEN 0.17 Non-Reactive: <1.0 **ECLIA**

> Reactive: =/>1.0 Source: Roche IFU.

INTERPRETATION NOTES:

1. A negative test result does not completely rule out the possibility of an infection with HIV. Serum or plasma samples from the very early (preseroconversion) phase or the late phase of HIV infection can occasionally yield negative findings. Yet unknown HIV variants can also lead to a negative HIV finding. The presence of antibodies to HIV is not a diagnosis of AIDS.

2. For diagnostic purposes, the results should always be assessed in conjunction with the patient's medical history, clinical examination and other

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Dubai, UAE





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IMMUNOLOGY

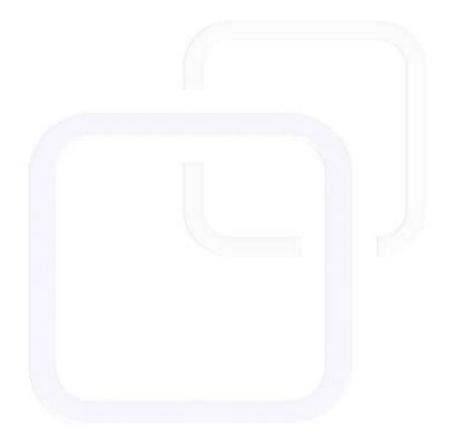
Test Result Flag Unit **Reference Range** Methodology

findings.

Name

3. This is a screening test.

Source: Roche Cobas IFU.



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SEROLOGY

TestResultFlagUnitReference RangeMethodologyRPR (RAPID PLASMA REAGEN)Non-reactiveNon-reactiveCarbon flocculation

INTERPRETATION NOTES:

Referred by

Syphilis is a disease caused by infection with the spirochete Treponema pallidum. The infection is systemic and the disease is characterized by periods of latency.

Patients with primary or secondary syphilis should be reexamined clinically and serologically 6 months and 12 months following treatment. Typically, rapid plasma reagin (RPR) titers decrease following successful treatment, but this may occur over a period of months to years. Biological false-positive reactions with cardiolipin-type antigens have been reported in disease such as infectious mononucleosis, leprosy, malaria, lupus erythematosus, vaccinia, and viral pneumonia. Pregnancy, autoimmune diseases, and narcotic addictions may give false-positives. Pinta, yaws, bejel, and other treponemal diseases may also produce false-positive results with this test.

False negatives tend to be more common in the initial and end stages of infection. Among people who are in the secondary (middle) stage of infection, the RPR test result is nearly always positive. (Interpretation added on 28 Dec 2019).

Sample Type : Serum

End of Report

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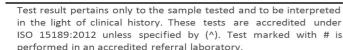
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