

Al Abbar Laboratories for Research and Medical Analysis مختبرات العبارللأبحاث و التحاليل الطبية



إدارة دبن للاعتماد طبقا لمواصفات الأيزو ١٥١٨٩ 15189 Dubai Accreditation Department for ISO 15189

LABORATORY REPORT

Name : PEN EI PHYU File. No. : AAL02-360665

DOB/Gender

: 26-05-1995 (28 Yrs 5 Month 28 Days/Female) Referral Doctor

: Dr. Sajid Sanaullah Khan

: 22233270305

Lab No.

Referral Clinic

: Peshawar(Irham Medical Center)

Request Date : 23-11-2023 21:05

Insurance : No Clinic File No : 41561

IMMUNOLOGY/SEROLOGY/INFECTIOUS DISEASES

Test Name	Result	Units	Ref. Range	Method
**Hepatitis B surface Ag	0.40	Cutoff Index (COI)	Non Reactive: <0.90 Borderline: >= 0.90 to < 1.0 Reactive: >= 1.0	ECLIA
**Hepatitis B "s" Ag Interpretation	Non-Reactive		Non-Reactive	

Full HepB profile is recommended in case of a Reactive result. HBs Ag Index value is an instrument dependant value and should not be used for monitoring.

For monitoring HepB levels (track treatment), a quantitative viral load by PCR is advised.

Samples should not be taken from patients receiving therapy with high biotin doses (i.e. > 5 mg/day) until at least 8 hours following the last biotin administration.

Please note Reference Range Reviewed w.ef 20/10/23

*HCV Abs (serum)

S/CO

Non-Reactive: < 1 Reactive: >= 1

CMIA

**Hepatitis C Virus Ab Interpretation

Non-Reactive

Non-Reactive

Note: Patient with acute HCV infection may still be non-reactive for HCV abs ('diagnostic window'). Samples should not be taken from patients receiving therapy with high biotin doses (i.e. > 5 mg/day) until at least 8 hours following the last biotin administration.

Please note Reference Range Reviewed w.ef 15/10/23

**HIV I & II Abs + p24 Ag

Cutoff Index (COI)

Non Reactive: <1.0 Reactive: >=1.0

ECLIA

**HIV Interpretation

Non-Reactive

Non-Reactive

Kindly note above test is just a screening test.

Note: Patients at a very early stage of infection may still be non-reactive for HIV Abs or detect p24 antigen. If HIV is suspected, a control sample is recommended in 4-6 weeks.

Samples should not be taken from patients receiving therapy with high biotin doses (i.e. > 5 mg/day) until at least 8 hours following the last biotin administration.

Please note Reference Range Reviewed w.ef 15/10/23

Sample Type: Serum

----- End Of Report ------

These tests are accredited under ISO 15189:2012 unless specified by (*) Sample processed on the same day of receipt unless specified otherwise. Test results pertains only the sample tested and to be correlated with clinical history.

Reference range related to Age/Gender.

Dr. Solmaz Siddiqui

Authenticated On: 23-11-2023 23:12:48 Printed On: 23-11-2023 23:21:46

H/L Collected On: 23-11-2023 19:00:00 Received On: 23-11-2023 21:06:00

Reprinted On: 24-11-2023 09:34:26

Released On: 23-11-2023 23:19:18

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