

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



إدارة دبن للإعتماد طبقا لمواصفات الأيزو ١٥١٨٩ ما المام Dubai Accreditation Department for ISO 15189

LABORATORY REPORT

Name : UTTARAM GURUNG KANCHHA GURUNG File. No. : AAL02-429255

DOB/Gender : 16-11-2000 (24 Yrs 1 Month 1 Days/Male) Referral Doctor : Dr. Humaira Mumtaz

Lab No. : 22243510178 Referral Clinic : Citicare Medical Center LLC

Insurance : NATIONAL GENERAL INSURANCE (HEALTHNET)

HAEMATOLOGY & COAGULATION

Test Name	Result	Units	Ref. Range	Method
COMPLETE BLOOD COUNT (C	BC) WITH DIFFERENTI	AL		
RBC	4.85	10^12/L	4.50 - 5.50	Hydrodynamic focusing (DC Detection)
Haemoglobin	14.3	g/dl	13.0 - 17.0	Photometry-SLS
нст	42.2	%	40.0 - 50.0	Hydrodynamic focusing (HF)
MCV	87.0	fl	83.0 - 101.0	Calculation
MCH	29.5	pg	27-32	Calculation
MCHC	33.9	g/dL	31.5 - 34.5	Calculation
Platelet Count	289	10^3/uL	150 - 400	HF (DCD)
WBC	10.21 ^H	10^3/uL	4.00 - 10.00	Flow Cytometry
DIFFERENTIAL COUNT (%)				
Neutrophils	61.9	%	40.0 - 80.0	Flow Cytometry
Lymphocytes	28.5	%	20.0 - 40.0	Flow Cytometry
Monocytes	5.3	%	2.0-10.0	Flow Cytometry
Eosinophils	3.3	%	1 - 6	Flow Cytometry
Basophils	1.0	%	<1-2	Flow Cytometry
Band Forms	0.0	%	< 6	Flow Cytometry
DIFFERENTIAL COUNT (ABSO	LUTE)			
Neutrophils (Absolute)	6.32	10^3/uL	2.00 - 7.00	Calculation
Lymphocytes (Absolute)	2.91	10^3/uL	1.00 - 3.00	Calculation
Monocytes (Absolute)	0.54	10^3/uL	0.20 - 1.00	Calculation
Eosinophils (Absolute)	0.34	10^3/uL	0.02 - 0.50	Calculation
Basophils (Absolute)	0.10	10^3/uL	0.02 - 0.10	Calculation
Band Forms (Absolute)	0.00	10^3/uL	< 0.66	Calculation

Remarks:

Test result to be interpreted in the light of clinical history and to be investigated further if necessary.

Sample Type : EDTA WB

These tests are accredited under ISO 15189:2022 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

Laboratory Director DHA/LS/248469 Patient Sample Collected On: 15-12-2024 05:53:00 Received On: 16-12-2024 14:21:00

Authenticated On: 16-12-2024 15:54:26 Released On: 16-12-2024 16:51:28

eleased On : 16-12-2024 16:51:28 **Kur** Sr. ٦

Pings

Kumareshan Sr. Technician DHA-P-2992011

Padmapriya





DOB/Gender

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



: AAL02-429255

: Dr. Humaira Mumtaz

: Citicare Medical Center LLC

إدارة دبى للإعتماد طبقا لمواصفات الأيزو ١٥١٨٩ - Dubai Accreditation Department for ISO 15189

LABORATORY REPORT

: UTTARAM GURUNG KANCHHA GURUNG Name

File. No. : 16-11-2000 (24 Yrs 1 Month 1 Days/Male) Referral Doctor

Lab No. : 22243510178 Referral Clinic

Request Date : 16-12-2024 14:19:32 Clinic File No : 43886

: NATIONAL GENERAL INSURANCE (HEALTHNET) Insurance

CLINICAL BIOCHEMISTRY

Test Name	Result	Units	Ref. Range	Method
C-Reactive Protein (CRP)	<0.60	mg/L	Negative < or = to 5.0	Immunoturbidim etric Assay

CRP is an acute phase protein whose concentration rises non -specifically in response to inflammation. CRP values should not be interpreted without a complete clinical evaluation. Follow-up testing of patients with elevated values is recommended in order to help rule out a recent response to undetected infection or tissue injury. Please note Reference Range Reviewed w.ef 15/10/23

Sample Type : Serum

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Dr. Solmaz Siddiqui Laboratory Director DHA/LS/248469

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Padmapriya Kumareshan

Sr. Technician DHA-P-2992011





DOB/Gender

Insurance

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LABORATORY REPORT

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: 16-11-2000 (24 Yrs 1 Month 1 Days/Male)

Lab No. : **22243510178**

Request Date : 16-12-2024 14:19:32

Referral Doctor Referral Clinic

File. No.

: Dr. Humaira Mumtaz

: AAL02-429255

: Citicare Medical Center LLC

Clinic File No : 43886

: NATIONAL GENERAL INSURANCE (HEALTHNET)

IMMUNOLOGY/SEROLOGY/INFECTIOUS DISEASES

Test Name	Result	Units	Ref. Range	Method
H.pylori Abs - IgG	32.8 ^H	AU/mL	Negative: < 8 Doubtfull: 8 - 12	ELISA

Positive: > 12

A positive test result confirms the presence of specific antibodies.

A negative result indicates that no clinically relevant antibodies against the pathogen are present in the patient's sample, but does not exclude the possibility of an acute infection.

In case of a borderline result a reliable evaluation is not possible.

A definitive diagnosis can only be achieved by testing paired serum samples, taken at one to two weeks intervals, in parallel.

Remarks:

Test result to be interpreted in the light of clinical history and to be investigated further if necessary.

Sample Type : Serum

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

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Dr. Solmaz SiddiquiLaboratory Director

DHA/LS/248469

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Padmapriya Kumareshan

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