

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 : AMEEN HOUSSEN ALKOURDI

DOB/Gender : 10-10-1983 (40 Yrs 11 Month 21 Days/Male)

Lab No. : **22242440097**

Request Date : 31-08-2024 13:22

Insurance : NEXTCARE

File. No. : **DML02-288715**

Referral Doctor : Dr. Enomen Goodluck Ekata
Referral Clinic : Citicare Medical Center LLC

Clinic File No : 38201

HAEMATOLOGY & COAGULATION

Test Name	Result	Units	Ref. Range	Method
COMPLETE BLOOD COUNT (CE	BC) WITH DIFFERENT	<u>IAL</u>		
RBC	5.62 ^H	10^12/L	4.50 - 5.50	Hydrodynamic focusing (DC Detection)
Haemoglobin	15.3	g/dl	13.0 - 17.0	Photometry-SLS
НСТ	45.7	%	40.0 - 50.0	Hydrodynamic focusing (HF)
MCV	81.3 ^L	fl	83.0 - 101.0	Calculation
MCH	27.2	pg	27-32	Calculation
MCHC	33.5	g/dL	31.5 - 34.5	Calculation
Platelet Count	201	10^3/uL	150 - 400	HF (DCD)
WBC	6.80	10^3/uL	4.00 - 10.00	Flow Cytometry
DIFFERENTIAL COUNT (%)				
Neutrophils	49.4	%	40.0 - 80.0	Flow Cytometry
Lymphocytes	38.4	%	20.0 - 40.0	Flow Cytometry
Monocytes	9.0	%	2.0-10.0	Flow Cytometry
Eosinophils	2.5	%	1 - 6	Flow Cytometry
Basophils	0.7	%	<1-2	Flow Cytometry
Band Forms	0.0	%	< 6	Flow Cytometry
DIFFERENTIAL COUNT (ABSOL	.UTE)			
Neutrophils (Absolute)	3.36	10^3/uL	2.00 - 7.00	Calculation
Lymphocytes (Absolute)	2.61	10^3/uL	1.00 - 3.00	Calculation
Monocytes (Absolute)	0.61	10^3/uL	0.20 - 1.00	Calculation
Eosinophils (Absolute)	0.17	10^3/uL	0.02 - 0.50	Calculation
Basophils (Absolute)	0.05	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 15189:2012 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 SiddiquiLaboratory Director

DHA/LS/248469

Patient Sample Collected On: 31-08-2024 12:30:00 Authenticated On: 31-08-2024 14:15:45

Printed On: 31-08-2024 18:55:58

Received On: 31-08-2024 13:27:00

Released On: 31-08-2024 15:22:06

:22:06 Laboratory Technician DHA-P-0039673





Wasik Hasan Tisekar



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



FCLIA

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Clinic File No : 38201

CLINICAL BIOCHEMISTRY

Test Name Result Units Ref. Range Method **C-Reactive Protein (CRP)** 2.58 Negative < or = to 5.0 Immunoturbidim mg/L 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

Troponin T (Quantitative) 0.0042

< 0.014 : Healthy Individuals

Risk Stratification in Asymptomatic Individuals <0.005 : Low Risk

>=0.005 to <=0.010 : Moderate

>0.010: High

Conversion Formula: (Concentration in pg/mL) x (0.001) = ng/mL. Values should be correlated with the clinical findings and ECG picture along with CKMB (Mass)and Myoglobin levels, before ruling out AMI. A sequential follow-up of Troponin T levels is recommended 1 hours post onset of symptoms / post admission for suspected AMI cases. Diagnostic criteria for AMI is defined as > 0.014 ng/ml. Please note Reference Range Reviewed w.ef 15/10/23 Sample Type: Serum

--- End Of Report ---

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Dr. Solmaz Siddiqui Laboratory Director DHA/LS/248469

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