

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 : MUHAMMAD NAWAZ MUHAMMAD IRFAN

DOB/Gender : 01-01-1983 (40 Yrs 11 Month 29 Days/Male)

Lab No. : **22233640351**

Request Date : 30-12-2023 21:17:59

Insurance : No

File. No. : AAL02-368104

Referral Doctor : **Dr. Enomen Goodluck Ekata**Referral Clinic : **Peshawar(Irham Medical Center)**

Clinic File No : 27703

HAEMATOLOGY & COAGULATION

Test Name	Result	Units	Ref. Range	Method
COMPLETE BLOOD COUNT (CBC)	WITH DIFFEREN	TIAL		
RBC	5.10	10^12/L	4.50 - 5.50	Hydrodynamic focusing (DC Detection)
Haemoglobin	13.8	g/dl	13.0 - 17.0	Photometry-SLS
НСТ	41.0	%	40.0 - 50.0	Hydrodynamic focusing (HF)
MCV	80.4 ^L	fl	83.0 - 101.0	Calculation
MCH	27.1	pg	27-32	Calculation
MCHC	33.7	g/dL	31.5 - 34.5	Calculation
Platelet Count	264	10^3/uL	150 - 400	HF (DCD)
WBC	7.72	10^3/uL	4.00 - 10.00	Flow Cytometry
DIFFERENTIAL COUNT (%)				
Neutrophils	58.7	%	40.0 - 80.0	Flow Cytometry
Lymphocytes	22.0	%	20.0 - 40.0	Flow Cytometry
Monocytes	9.8	%	2.0-10.0	Flow Cytometry
Eosinophils	9.1 ^H	%	1 - 6	Flow Cytometry
Basophils	0.4	%	<1-2	Flow Cytometry
Band Forms	0.0	%	< 6	Flow Cytometry
DIFFERENTIAL COUNT (ABSOLUTE	<u>5</u>)			
Neutrophils (Absolute)	4.53	10^3/uL	2.00 - 7.00	Calculation
Lymphocytes (Absolute)	1.70	10^3/uL	1.00 - 3.00	Calculation
Monocytes (Absolute)	0.76	10^3/uL	0.20 - 1.00	Calculation
Eosinophils (Absolute)	0.70 ^H	10^3/uL	0.02 - 0.50	Calculation
Basophils (Absolute)	0.03	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 Siddiqui Laboratory Director DHA/LS/248469 H/L Collected On: 30-12-2023 20:00:00 Received On: 30-12-2023 21:18:00
Authenticated On: 30-12-2023 23:59:03 Released On: 31-12-2023 00:05:21

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Esmeralda Obenita Lab Incharge DHA/LS/2992011/ 241430





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Referral Clinic : Peshawar(Irham Medical Center)

Clinic File No : 27703

CLINICAL BIOCHEMISTRY

Test Name	Result	Units	Ref. Range	Method
C-Reactive Protein (CRP)	7.95 ^H	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 Remarks:

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

Sample Type : Serum

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ECLIA

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LABORATORY REPORT

Name : MUHAMMAD NAWAZ MUHAMMAD IRFAN

DOB/Gender : 01-01-1983 (40 Yrs 11 Month 29 Days/Male)

Lab No. : 22233640351

Request Date : 30-12-2023 21:17:59

Insurance : No File. No. : AAL02-368104

Referral Doctor : Dr. Enomen Goodluck Ekata

Referral Clinic : Peshawar(Irham Medical Center)

Clinic File No : 27703

IMMUNOLOGY/SEROLOGY/INFECTIOUS DISEASES

Test Name Result Units Ref. Range Method

Non Reactive: <0.90 **Hepatitis B surface Ag 0.39 Cutoff Index (COI)

Borderline: >= 0.90 to < 1.0 Reactive: >= 1.0

**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

Hepatitis C Virus Abs ECLIA Cutoff Index (COI) Non Reactive: < 0.90

Borderline: >= 0.90 to < 1.0

Reactive: >=1.0

Non-Reactive **Hepatitis C Virus Ab Interpretation** 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

Sample Type : Serum

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Dr. Solmaz Siddiqui Laboratory Director

DHA/LS/248469

Received On: 30-12-2023 21:28:00 H/L Collected On: 30-12-2023 20:00:00 Authenticated On: 30-12-2023 23:59:04

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CLINICAL BIOCHEMISTRY						
Test Name	Result	Units	Ref. Range	Method		
Bilirubin - Total	0.26	mg/dL	0.20-1.20	DIAZO		
Please note Reference Range Reviewed w.	ef 15/10/23					
Bilirubin - Direct	0.16	mg/dL	0.00-0.5	DIAZO		
Please note Reference Range Reviewed w.	ef 15/10/23					
Alanine Aminotransferase (ALT / GPT)	47.90	U/L	10.0 - 50.0	NADH w/ P5P		
Please note Reference Range Reviewed w.	ef 15/10/23					
Aspartate Aminotransferase(AST / GOT)	27.80	U/L	10 - 50	NADH w/ P5P		
Please note Reference Range Reviewed w.	ef 15/10/23					
Alkaline Phosphatase	101.00	U/L	40-129	Colorimetric -P- NPT		
Please note Reference Range Reviewed w.						
Gamma Glutamyl Transferase (GGT)) 96.30 ^H	U/L	<60	Enzymatic-GG 3C4Nitro AN		
Please note Reference Range Reviewed w.	ef 15/10/23					
Protein - Total	6.54	g/dL	6.40 - 8.30	Colorimetric - Biuret		
Please note Reference Range Reviewed w.	ef 15/10/23.					
Albumin	4.08	g/dL	3.97 - 4.94	Colorimetric - BCG		
Please note Reference Range Reviewed w.	ef 15/10/23					
*Globulin (serum)	2.46 ^L	g/dL	2.50 - 3.50	Calculation		
**Albumin/Globulin Ratio	1.66	Ratio	1.10 - 2.50	Calculation		
Remarks:						
Test result to be interpreted in the light	of clinical history a	nd to be investigated f	urther if necessary.			

Sample Type: Serum

--- End Of Report -----

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