

LABORATORY REPORT

Name	: NASHWA FOUAD ARTIN	File. No.	: AAL02-425317
DOB/Gender	: 21-09-1973 (51 Yrs 5 Month 18 Days/Female)	Referral Doctor	: Dr. Humaira Mumtaz
Lab No.	: 22250370401	Referral Clinic	: Citicare Medical Center LLC
Request Date	: 06-02-2025 21:55:44	Clinic File No	: 44900
Insurance	: ALMADALLAH HEALTHCARE MANAGEMENT		

CLINICAL BIOCHEMISTRY

Test Name	Result	Units	Ref. Range	Method
Uric Acid	4.23	mg/dL	2.40 - 5.70	Uricase

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

Sample Type : Serum

These tests are accredited under ISO 15189:2022 unless specified by (*)

Sample processed on the same day of receipt unless specified otherwise.

Test results pertain 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 : 06-02-2025 18:27:00
Authenticated On : 06-02-2025 23:39:14

Received On : 06-02-2025 21:55:00
Released On : 06-02-2025 23:39:49



Esmeralda Obenita
Lab Incharge
DHA-P-2992011

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**LIPID PROFILE I
CLINICAL BIOCHEMISTRY**

Test Name	Result	Units	Ref. Range	Method
Cholesterol	240.00	mg/dL	Desirable: < 200 Borderline: 200 - 239 High: >or =240	Enzymatic- Cholesterol esterase
Please note Reference Range Reviewed w.ef 15/10/23				
Triglycerides	173.00 ^H	mg/dL	NCEP Adult Treatment Panel Normal: <150 Borderline: 150 - 199 High: 200-499 Very High: >or =500	Enzymatic GPO
Please note Reference Range Reviewed w.ef 15/10/23				
HDL Cholesterol	68.30	mg/dL	Major Risk: < 40	Homogeneous enzymatic-CHER & CHOD
Please note Reference Range Reviewed w.ef 15/10/23				
LDL Cholesterol	152.00 ^H	mg/dL	Optimal: < 100 Near or above optimal: 100-129 Borderline High: 130 - 159 High: 160 - 189 Very High: >or=190	Homo. Enz, Liq. Selective Surfactant
Please note Reference Range Reviewed w.ef 15/10/23				
*VLDL	19.70	mg/dL	5.0 - 30.0	Calculation
*Cholesterol / HDL	3.51	Ratio	Desirable: < 4.5 Borderline: 4.5 - 6 Undesirable: >6	Calculation

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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