

LABORATORY REPORT

Name	: SANDI AL NIMRI	File. No.	: AAL02-1307008
DOB/Gender	: 23-11-1989 (35 Yrs 5 Month 2 Days/Female)	Referral Doctor	: Dr. Aisha Umer
Lab No.	: 22251150277	Referral Clinic	: Citicare Medical Center LLC
Request Date	: 25-04-2025 21:04:00	Clinic File No	: 46497
Insurance	: NEXTCARE		

IMMUNOLOGY/SEROLOGY/INFECTIOUS DISEASES

Test Name	Result	Units	Ref. Range	Method
ATPO (Thyroid Peroxidase Antibodies)	38.40^H	IU/mL	< 34 Pregnancy: 1st Trimester: < 64 2nd Trimester: < 51 3rd Trimester: < 123	ECLIA

Anti-TPO antibodies are found often in conjunction with anti-Tg in the majority of cases of Hashimoto's thyroiditis, Primary Myxedema, and Graves' disease. Anti-TPO antibodies are demonstrable in most cases of postpartum thyroiditis and it has been found that the presence of autoantibody in early pregnancy is associated with a high risk of asymptomatic postpartum hypothyroidism. 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

Remarks:

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

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 : 25-04-2025 14:20:00
Authenticated On : 25-04-2025 22:35:11

Received On : 25-04-2025 21:10:00
Released On : 25-04-2025 22:42:00

Pring

Padmapriya Kumaresan
Sr. Technician
DHA-P-2992011

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HORMONES/ENDOCRINE/TUMOR MARKERS

Test Name	Result	Units	Ref. Range	Method
Prolactin	12.60	ng/mL	Non Pregnant: 4.79-23.3 Pregnant: 95-473	ECLIA

Conversion Formula: (Concentration in ng/mL) x (21.2) = mIU/L.

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 10/01/24

Sample Type : Serum

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

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

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Test results pertains only the sample tested and to be correlated with clinical history.

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