

Tests you can trust

Date : XX XX XXXX

Test Asked: Amh

Report Status: Complete Report



9 out of 10 Doctors trust that Thyrocare reports are accurate & reliable*







700+ Tests & Profiles



Temperature-Controlled Sample Logistics



Unique Barcode Tracking & Reports with QR Code Verification



Fully Automated Machines Inspected Daily



Abnormal Values Re-Checked Twice



Reports Verified By Expert MD Pathologists Stationed at Every Lab

Accredited by





ISO 9001: 2015 - From 2015



CAP From 2007

PROCESSED AT:





NAME : XXXXXXXXXXXXXXXX **SAMPLE COLLECTED AT:**

REF. BY

TEST ASKED : AMH

Report Availability Summary

Note: Please refer to the table below for status of your tests.

1 Ready

O Ready with Cancellation (Processing

TEST DETAILS REPORT STATUS

ANTI MULLERIAN HORMONE (AMH)



PROCESSED AT:





🗣 Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400703 🙎 98706 66333 🎽 wellness@thyrocare.com



9 out of 10 Doctors Trust that Thyrocare Reports are Accurate & Reliable

SAMPLE COLLECTED AT: NAME

REF. BY : XXXXXXXXXXXXXXXXX

: AMH **TEST ASKED**

PATIENTID : XXXXXXXXXXXXXXXXX

VALUE TEST NAME TECHNOLOGY UNITS ANTI MULLERIAN HORMONE (AMH) E.C.L.I.A 0.02 ng/mL

Bio. Ref. Interval. :-

Healthy Male: 0.77-14.5

Females

<= 2 years : <4.7 02-12 years: <8.8 12-20 years: 0.9-9.5 20-24 years :1.22-11.7 25-29 years: 0.89-9.85 30-34 years: 0.57-8.13 35-39 years: 0.14-7.49 40-44 years: 0.02-5.47 45-50 years: 0.010-2.71

PCOS women: 1.86-18.9

Clinical Significance:

- 1. Antimullerian hormone (AMH), also known as mullerian inhibiting substance, belongs to the transforming growth factor-beta
- 2. It is produced by Sertoli cells of the testis in males and by ovarian granulosa cells in females.
- 3. AMH is expressed in the follicles of females of reproductive age and inhibits the transition of follicles from primordial to primary
- 4. Measurement of AMH has utility in the assessment of gender, gonadal function, fertility, and as a gonadal tumor marker.
- 5. Since AMH is produced continuously in the granulosa cells of small follicles during the menstrual cycles, it is superior to the episodically released gonadotropins and ovarian steroids as a marker of ovarian reserve.
- 6. The monoclonal antibodies used are highly specific to Human AMH but it may cross react with high levels of LH, FSH, Inhibin A, Activin A, biotin etc.

References:

- 1. Kit Inserts
- 2. Wilson CA, di Clemente N, Ehrenfels C, et al. Mullerian inhibiting substance requires its N-terminal domain for maintenance of biological activity, a novel finding within the transforming growth factor-beta superfamily. Mol Endocrinol 1993;7(2):247-257.

Please correlate with clinical conditions.

Method:-SANDWICH ELECTRO-CHEMILUMINESCENCE IMMUNOASSAY (E.C.L.I.A)

~~ End of report ~~

Sample Collected on (SCT)

: Sample collection time

Sample Received on (SRT)

: Sample receiving time at Lab

Report Released on (RRT)

: Report release time

Sample Type

: SERUM

Doctor 1 Sign Doctor 2 Sign

Labcode

:

Barcode Page: 1 of 2 :

CONDITIONS OF REPORTING

- v The reported results are for information and interpretation of the referring doctor only.
- v It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- v Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for the same patient.
- v Should the results indicate an unexpected abnormality, the same should be reconfirmed.
- v Only such medical professionals who understand reporting units, reference ranges and limitations of technologies should interpret results.
- v This report is not valid for medico-legal purpose.
- v Neither Thyrocare, nor its employees/representatives assume any liability, responsibility for any loss or damage that may be incurred by any person as a result of presuming the meaning or contents of the report.
- v Thyrocare Discovery video link :- https://youtu.be/nbdYeRqYyQc
- v For clinical support please contact @8450950852,8450950853,8450950854 between 10:00 to 18:00

EXPLANATIONS

- v Majority of the specimen processed in the laboratory are collected by Pathologists and Hospitals we call them as "Clients".
- v Name The name is as declared by the client and recored by the personnel who collected the specimen.
- v Ref.Dr The name of the doctor who has recommended testing as declared by the client.
- v Labcode This is the accession number in our laboratory and it helps us in archiving and retrieving the data.
- v **Barcode** This is the specimen identity number and it states that the results are for the specimen bearing the barcode (irrespective of the name).
- v SCP Specimen Collection Point This is the location where the blood or specimen was collected as declared by the client.
- v SCT Specimen Collection Time The time when specimen was collected as declared by the client.
- v SRT Specimen Receiving Time This time when the specimen reached our laboratory.
- v RRT Report Releasing Time The time when our pathologist has released the values for Reporting.
- v Reference Range Means the range of values in which 95% of the normal population would fall.

SUGGESTIONS

- v Values out of reference range requires reconfirmation before starting any medical treatment.
- v Retesting is needed if you suspect any quality shortcomings.
- v Testing or retesting should be done in accredited laboratories.
- v For suggestions, complaints or feedback, write to us at info@thyrocare.com or call us on 022-3090 0000 / 62 360
- v SMS:<Labcode No.≯to 99666333

