PROCESSED AT :

Thyrocare

Thyrocare
Tests you can trust

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

NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : HE4

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
HUMAN EPIDIDYMIS PROTEIN 4	C.M.I.A	45.3	pmol/L

Bio. Ref. Interval. :-

Females :
Pre Menopausal : <=70 pmol/L
Post Menopausal : <=140 pmol/L

Interpretation:-
The level of HE4 cannot be used as an absolute evidence for presence or absence of malignant disease and HE4 assay should not be used as a cancer screening test.
The results obtained should be interpreted in conjunction with other methods in accordance with standard clinical management guidelines. If HE4 results are inconsistent with clinical evidence, additional testing is recommended.
Patients with confirmed ovarian cancer may have HE4 values similar to normal healthy women. The use of this assay is not recommended in monitoring patients with known mucinous or germ cell tumors (as these rarely produce HE4) . Elevated levels of HE4 antigen may be seen in patients with non malignant disease.
Specimens containing HAMA (Human anti mouse antibodies), heterophile antibodies etc may give anomalous values . The values may show variation if different methodologies, technologies, reagent brand, calibrators used and also variation seen due to various pathological and physiological factors

Reference - Abbott HE4 Reagent Kit insert

Please correlate with clinical conditions.

Method:- FULLY AUTOMATED CHEMI LUMINESCENT MICROPARTICLE IMMUNOASSAY

~~ End of report ~~

Dummy 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
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

CONDITIONS OF REPORTING

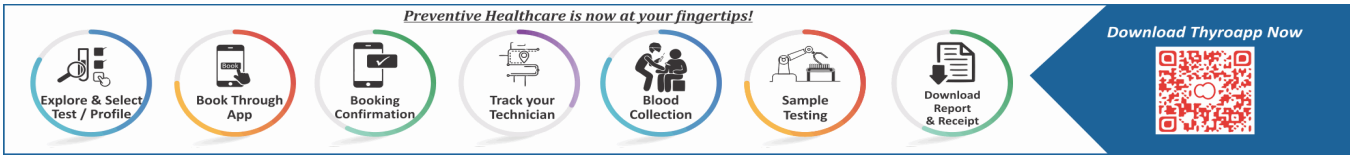
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for the same patient.
- Should the results indicate an unexpected abnormality, the same should be reconfirmed.
- Only such medical professionals who understand reporting units, reference ranges and limitations of technologies should interpret results.
- This report is not valid for medico-legal purpose.
- 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.
- Thyrocare Discovery video link :- https://youtu.be/nbdYeRgYyQc
- For clinical support please contact @8450950852,8450950853,8450950854 between 10:00 to 18:00

EXPLANATIONS

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

SUGGESTIONS

- Values out of reference range requires reconfirmation before starting any medical treatment.
- Retesting is needed if you suspect any quality shortcomings.
- Testing or retesting should be done in accredited laboratories.
- For suggestions, complaints or feedback, write to us at **info@thyrocare.com** or call us on **022-3090 0000 / 67 30 00**
- SMS:<Labcode No.>to **9[illegible]666333**





**As per a survey on doctors' perception of laboratory diagnostics (IJARIIT,2023)*