

Patient Name Centre
Age/Gender OP/IP No

Max ID/Mobile Collection Date/Time
Lab ID Receiving Date
Ref Doctor Reporting Date

Passport No.

Immunoassay

Max-Fertility hormonal profile

Test Name Result Unit Bio Ref Interval

TSH Camp*

Chemiluminescence

TSH 2.61 ulU/ml 0.34-5.6

Anti Mullerian Hormone (AMH)*

Anti Mullerian Hormone (AMH) **0.11** ng/mL 0.96 - 13.34

CLIA

Ref Range Interpretation:

Anti-Mullerian Hormone (AMH) is a hormone secreted by cells in developing egg sacs (follicles). The level of AMH in blood is generally a good indicator of ovarian reserve.

Low AMH levels are considered to be an indicator of a **low ovarian reserve**, i.e. few remaining follicles. AMH levels decline with age, and in younger women this may be a sign of premature loss of fertility

AMH does not change during menstrual cycle, so the blood sample can be taken at any time of the month - even while using oral contraception. AMH level for a fertile woman is 1.0–4.0 ng/ml

In males AMH is secreted by immature Sertoli cells (SC) and is responsible for the regression of Müllerian ducts in the male fetus as part of the sexual differentiation process. AMH is also involved in testicular development and function.

AMH level ng/ml	Effects for fertility treatment	
<0.4	Very low value. Very few eggs at stimulation. Pregnancy chances significantly low.	
0.4 - 1.0	Low value. Treatment may be possible.	
1.0 – 3.5	Normal value. Good possibility of treatment.	
>3.5	Suggestive of ovarian hyperstimulation syndrome / PCOS	

Note:- Optimal ovarian reserve values range between 2 - 6 ng/mL in reproductive age group



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SIN No:B2B1557895, Test Performed at :585 - Max Hospital - Gurugram, Opposite HUDA City Centre Metro Station, B - Block Booking Centre :3284 - Max Lab Sector 56 Gurgaon, House No 154, Ground Floor Near Kendriya Vihar, 9643211955 The authenticity of the report can be verified by scanning the Q R Code on top of the page

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Max Super Speciality Hospital, Saket (West Block), 1, Press Enclave Road, Saket, New Delhi - 110 017, Phone: +91-11-6611 5050 (CIN No.: U85100DL2021PLC381826)

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Estradiol (E2),Serum

Date 13/Mar/2022 Unit Bio Ref 12:31AM Interval

Estradiol <20 pg/mL

Ref Range

Passport No.

Male	20 - 75	
Post - Menopausal (Female)	20 - 88	
Estradiol -Total (Non - Pregnant Females)		
Mid Follicular Phase	24 - 114	
Mid - Luteal Phase	80 - 273	
Periovulatory	62 - 534	



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FSH - Follicle Stimulating Hormone, Serum*

Date 13/Mar/2022 Unit Bio Ref 12:31AM Interval

Follicle Stimulating 43.70 mlU/mL

Hormone

Ref. Range

Passport No.

Interpretation

Increased in primary gonadal failure, ovarian or testicular agenesis, Klinefelter's syndrome, Reifenstein's syndrome, castration, alcoholism, menopause, orchitis, gonadotropin-secreting pitutary tumors.

Decreased in anterior hypofunction, hypothalamic disorders, pregnancy, anorexia nervose, polycystic ovarian disease, hemochromatosis, sickle cell anaema, sever illness, hyperprolactinemia.

Pooled samples are advisable due to episodic, diurnal and cyclic variations in gonadotropin secretion.



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LH-Luteinizing Hormone, Serum*

Date 13/Mar/2022 Unit Bio Ref 12:31AM Interval

Luteinizing Hormone 7.86 mlU/mL

CLIA

Ref Range

LH(Male-Adult)	Reference Range	
	1.24-8.62	
LH (Female-Adult)		
Follicular	2.12-10.89	
Mid Cycle Peak	19.18-103.03	
Luteal Phase	1.2-12.86	
Post Menopausal (>50 Year)	10.87-58.64	

Interpretation

Increased in Primary gonadal dysfunction, polycystic ovarian syndrome (LH/FSH ratio is high in 60% cases), post-menopause, and pituitary adenoma. Decreased in pituitary or hypothalamic impairment, isolated gonadotropic deficiency associated with anosmia or hyposmia (Kallmann's syndrome), anorexia nervosa, isolated LH deficiency ("fertile eunuch"), sever stress, malnutrition, and sever illness.

Pooled samples are advisable due to episodic, diurnal and cyclic variations in gonadotropin secretion.

Prolactin, Serum*

Date 13/Mar/2022 Unit Bio Ref Interval

12:31AM

Prolactin 3.67 ng/mL

Ref Range

Males: 2.64 - 13.13

Females:
Premenopausal

(<50 years of 3.34 - 26.74

age):



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Postmenopausal

(>50 years of 2.74 - 19.64

age):

Interpretation

Increased in prolactin-secreting pituitary tumors, amenorrhea and/or galactorrhea, Chiari-Frommel and Argonz Del Cstillo syndromes, various types of hypothalamic-pitutary disease (e.g. sarcoidosis, granulomatous diseases, crangiopharyngioma, metastatic cancer, empty sella syndrome), primary hypothyroidism, anorexia nervosa, polycystic ovary syndrome, renal failure, insulin-induced hypoglycemia, chest wall injury, adrenal insufficiency, and pituitary stalk section surgery Decreased in pituitary apoplexy (Sheehan's Syndrome)

Kindly correlate with clinical findings

*** End Of Report ***

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