

Dr. Vinay Chopra  
 MD (Pathology & Microbiology)  
 Chairman & Consultant Pathologist

Dr. Yugam Chopra  
 MD (Pathology)  
 CEO & Consultant Pathologist

**NAME** : Mr. MANISH  
**AGE/ GENDER** : 46 YRS/MALE  
**COLLECTED BY** :  
**REFERRED BY** : DR. RESHAM SINGH  
**BARCODE NO.** : 01520384  
**CLIENT CODE.** : KOS DIAGNOSTIC LAB  
**CLIENT ADDRESS** : 6349/1, NICHOLSON ROAD, AMBALA CANTT

**PATIENT ID** : 1665620  
**REG. NO./LAB NO.** : 012411080054  
**REGISTRATION DATE** : 08/Nov/2024 03:10 PM  
**COLLECTION DATE** : 08/Nov/2024 03:10PM  
**REPORTING DATE** : 08/Nov/2024 05:48PM

| Test Name | Value | Unit | Biological Reference interval |
|-----------|-------|------|-------------------------------|
|-----------|-------|------|-------------------------------|

## ENDOCRINOLOGY

### THYROID FUNCTION TEST: TOTAL

|  |       |        |              |
|--|-------|--------|--------------|
| TRIIODOTHYRONINE (T3): SERUM<br>by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY)             | 0.938 | ng/mL  | 0.35 - 1.93  |
| THYROXINE (T4): SERUM<br>by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY)                    | 8.61  | µg/dL  | 4.87 - 12.60 |
| THYROID STIMULATING HORMONE (TSH): SERUM<br>by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY) | 1.755 | µIU/mL | 0.35 - 5.50  |

3rd GENERATION, ULTRASENSITIVE

#### INTERPRETATION:

TSH levels are subject to circadian variation, reaching peak levels between 2-4 a.m and at a minimum between 6-10 pm. The variation is of the order of 50%. Hence time of the day has influence on the measured serum TSH concentrations. TSH stimulates the production and secretion of the metabolically active hormones, thyroxine (T4) and triiodothyronine (T3). Failure at any level of regulation of the hypothalamic-pituitary-thyroid axis will result in either underproduction (hypothyroidism) or overproduction (hyperthyroidism) of T4 and/or T3.

| CLINICAL CONDITION           | T3                    | T4                    | TSH                             |
|------------------------------|-----------------------|-----------------------|---------------------------------|
| Primary Hypothyroidism:      | Reduced               | Reduced               | Increased (Significantly)       |
| Subclinical Hypothyroidism:  | Normal or Low Normal  | Normal or Low Normal  | High                            |
| Primary Hyperthyroidism:     | Increased             | Increased             | Reduced (at times undetectable) |
| Subclinical Hyperthyroidism: | Normal or High Normal | Normal or High Normal | Reduced                         |

#### LIMITATIONS:-

1. T3 and T4 circulates in reversibly bound form with Thyroid binding globulins (TBG), and to a lesser extent albumin and Thyroid binding Pre Albumin so conditions in which TBG and protein levels alter such as pregnancy, excess estrogens, androgens, anabolic steroids and glucocorticoids may falsely affect the T3 and T4 levels and may cause false thyroid values for thyroid function tests.
2. Normal levels of T4 can also be seen in Hyperthyroid patients with : T3 Thyrotoxicosis, Decreased binding capacity due to hypoproteinemia or ingestion of certain drugs (e.g.: phenytoin, salicylates).
3. Serum T4 levels in neonates and infants are higher than values in the normal adult, due to the increased concentration of TBG in neonate serum.
4. TSH may be normal in central hypothyroidism, recent rapid correction of hyperthyroidism or hypothyroidism, pregnancy, phenytoin therapy.

| TRIIODOTHYRONINE (T3) |                          | THYROXINE (T4)    |                          | THYROID STIMULATING HORMONE (TSH) |                          |
|-----------------------|--------------------------|-------------------|--------------------------|-----------------------------------|--------------------------|
| Age                   | Refferance Range (ng/mL) | Age               | Refferance Range (µg/dL) | Age                               | Reference Range (µIU/mL) |
| 0 - 7 Days            | 0.20 - 2.65              | 0 - 7 Days        | 5.90 - 18.58             | 0 - 7 Days                        | 2.43 - 24.3              |
| 7 Days - 3 Months     | 0.36 - 2.59              | 7 Days - 3 Months | 6.39 - 17.66             | 7 Days - 3 Months                 | 0.58 - 11.00             |
| 3 - 6 Months          | 0.51 - 2.52              | 3 - 6 Months      | 6.75 - 17.04             | 3 Days - 6 Months                 | 0.70 - 8.40              |
| 6 - 12 Months         | 0.74 - 2.40              | 6 - 12 Months     | 7.10 - 16.16             | 6 - 12 Months                     | 0.70 - 7.00              |



  
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| Test Name  | Value       | Unit                | Biological Reference interval |
|--|-------------|---------------------|-------------------------------|
| 1 - 10 Years   | 0.92 - 2.28 | 1 - 10 Years        | 6.00 - 13.80                  |
| 11- 19 Years   | 0.35 - 1.93 | 11 - 19 Years       | 4.87- 13.20                   |
| > 20 years (Adults)  | 0.35 - 1.93 | > 20 Years (Adults) | 4.87 - 12.60                  |
| RECOMMENDATIONS OF TSH LEVELS DURING PREGNANCY ( $\mu$ U/mL) |             |                     |                               |
| 1st Trimester  |             |                     | 0.10 - 2.50                   |
| 2nd Trimester  |             |                     | 0.20 - 3.00                   |
| 3rd Trimester  |             |                     | 0.30 - 4.10                   |

#### INCREASED TSH LEVELS:

- 1.Primary or untreated hypothyroidism may vary from 3 times to more than 100 times normal depending upon degree of hypofunction.
- 2.Hypothyroid patients receiving insufficient thyroid replacement therapy.
- 3.Hashimotos thyroiditis
- 4.DRUGS: Amphetamines, iodine containing agents & dopamine antagonist.
- 5.Neonatal period, increase in 1st 2-3 days of life due to post-natal surge

#### DECREASED TSH LEVELS:

- 1.Toxic multi-nodular goiter & Thyroiditis.
- 2.Over replacement of thyroid hormone in treatment of hypothyroidism.
- 3.Autonomously functioning Thyroid adenoma
- 4.Secondary pituitary or hypothalamic hypothyroidism
- 5.Acute psychiatric illness
- 6.Severe dehydration.
- 7.DRUGS: Glucocorticoids, Dopamine, Levodopa, T4 replacement therapy, Anti-thyroid drugs for thyrotoxicosis.
- 8.Pregnancy: 1st and 2nd Trimester



  
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### IMMUNOPATHOLOGY/SEROLOGY

#### HEPATITIS C VIRUS (HCV) ANTIBODY: TOTAL

|  |                |      |                  |
|--|----------------|------|------------------|
| HEPATITIS C ANTIBODY (HCV) TOTAL: SERUM              | 0.06           | S/CO | NEGATIVE: < 1.00 |
| by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY) |                |      | POSITIVE: > 1.00 |
| HEPATITIS C ANTIBODY (HCV) TOTAL RESULT              | NON - REACTIVE |      |                  |
| by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY) |                |      |                  |

#### INTERPRETATION:-

| RESULT (INDEX) | REMARKS  |
|----------------|--|
| < 1.00         | NON - REACTIVE/NOT - DETECTED                        |
| > =1.00        | REACTIVE/ASYMPTOMATIC/INFECTIVE STATE/CARRIER STATE. |

Hepatitis C (HCV) is an RNA virus of Favivirus group transmitted via blood transfusions, transplantation, injection drug abusers, accidental needle punctures in healthcare workers, dialysis patients and rarely from mother to infant. 10 % of new cases show sexual transmission. As compared to HAV & HBV , chronic infection with HCV occurs in 85 % of infected individuals. In high risk population, the predictive value of Anti HCV for HCV infection is > 99% whereas in low risk populations it is only 25 %.

#### USES:

- Indicator of past or present infection, but does not differentiate between Acute/ Chronic/Resolved Infection.
- Routine screening of low and high prevalence population including blood donors.

#### NOTE:

- False positive results are seen in Auto-immune disease, Rheumatoid Factor, HYpergammaglobulinemia, Paraproteinemia, Passive antibody transfer, Anti-idiotypes and Anti-superoxide dismutase.
- False negative results are seen in early Acute infection, Immunosuppression and Immuno— incompetence.
- HCV-RNA PCR recommended in all reactive results to differentiate between past and present infection.



  
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HEPATITIS B SURFACE ANTIGEN (HBsAg) ULTRA

HEPATITIS B SURFACE ANTIGEN (HBsAg): 0.34 S/CO  
SERUM  
by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY)  
NEGATIVE: < 1.0  
POSITIVE: > 1.0

HEPATITIS B SURFACE ANTIGEN (HBsAg) NON REACTIVE  
RESULT  
by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY)

INTERPRETATION:

| RESULT IN INDEX VALUE | REMARKS        |
|-----------------------|----------------|
| < 1.30                | NEGATIVE (-ve) |
| >=1.30                | POSITIVE (+ve) |

Hepatitis B Virus (HBV) is a member of the Hepadna virus family causing infection of the liver with extremely variable clinical features. Hepatitis B is transmitted primarily by body fluids especially serum and also spread effectively sexually and from mother to baby. In most individuals HBV hepatitis is self limiting, but 1-2 % normal adolescent and adults develop Chronic Hepatitis. Frequency of chronic HBV infection is 5-10% in immunocompromised patients and 80 % neonates. The initial serological marker of acute infection is HBsAg which typically appears 2-3 months after infection and disappears 12-20 weeks after onset of symptoms. Persistence of HBsAg for more than 6 months indicates carrier state or Chronic Liver disease.

\*\*\* End Of Report \*\*\*



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