



| | | Chopra gy & Microbiology) Consultant Pathologist | | (Pathology) |
|--|----------------------------------|---|--------------------------|--|
| NAME | : Mrs. ANJU SHARMA | | | |
| AGE/ GENDER | : 48 YRS/FEMALE | | PATIENT ID | : 1549160 |
| COLLECTED BY | : SURJESH | | REG. NO./LAB NO. | : 012407150038 |
| REFERRED BY | : | | REGISTRATION DATE | : 15/Jul/2024 10:26 AM |
| BARCODE NO. | :01513184 | | COLLECTION DATE | : 15/Jul/2024 10:31AM |
| CLIENT CODE. | : KOS DIAGNOSTIC LAB | | REPORTING DATE | : 15/Jul/2024 11:25AM |
| CLIENT ADDRESS | : 6349/1, NICHOLSON ROA | AD, AMBALA CANTT | | |
| Test Name | | Value | Unit | Biological Reference interval |
| | CL | INICAL CHEMIS | TRY/BIOCHEMISTR | v |
| | ŰĽ | | OFILE : BASIC | |
| CHOLESTEROL TOTAL | .: SERUM | 234.65 ^H | mg/dL | OPTIMAL: < 200.0 |
| by CHOLESTEROL OX | | 234.03 | g. all | BORDERLINE HIGH: 200.0 - 239.0 |
| | | | | HIGH CHOLESTEROL: > OR = 240. |
| FRIGLYCERIDES: SER | UIVI HATE OXIDASE (ENZYMATIC) | 222.87 ^H | mg/dL | OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 |
| | | | | HIGH: 200.0 - 499.0 |
| | | | | VERY HIGH: > OR = 500.0 |
| HDL CHOLESTEROL (DIRECT): SERUM by SELECTIVE INHIBITION | | 44.69 | mg/dL | LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 - |
| | | | | 60.0 |
| | | | | HIGH HDL: $> OR = 60.0$ |
| LDL CHOLESTEROL: S by CALCULATED, SPE | | 145.39 ^H | mg/dL | OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129.0 |
| · · · · · · · · · · · · · · · · · · · | | | | BORDERLINE HIGH: 130.0 - 159.0 |
| | | | | HIGH: 160.0 - 189.0 |
| | | 400 o/H | ma/dl | VERY HIGH: > OR = 190.0 OPTIMAL: < 130.0 |
| NON HDL CHOLESTEROL: SERUM by CALCULATED, SPECTROPHOTOMETRY | | 189.96 ^H | mg/dL | ABOVE OPTIMAL: 130.0 - 159.0 |
| | | | | BORDERLINE HIGH: 160.0 - 189.0 |
| | | | | HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0 |
| VLDL CHOLESTEROL: | | 44.57 | mg/dL | 0.00 - 45.00 |
| by CALCULATED, SPECTROPH | Λ | 692.17 | mg/dL | 350.00 - 700.00 |
| by CALCULATED, SPEC | | 5.25 ^H | RATIO | LOW RISK: 3.30 - 4.40 |
| by CALCULATED, SPE | | 5.25 | KATIU | AVERAGE RISK: 4.50 - 7.0 |
| | | | | MODERATE RISK: 7.10 - 11.0 |
| | | | | HIGH RISK: > 11.0 |
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KOS Diagnostic Lab (A Unit of KOS Healthcare)

DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY)

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DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST MBBS , MD (PATHOLOGY)

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Page 1 of 4





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| Test Name | | Value | Unit | Biological Reference interval |
| LDL/HDL RATIO: SEF by CALCULATED, SPI | | 3.25 ^H | RATIO | LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0 |
| TRIGLYCERIDES/HDL by CALCULATED, SPE | | 4.99 | RATIO | 3.00 - 5.00 |

INTERPRETATION:

1. Measurements in the same patient can show physiological analytical variations. Three serial samples 1 week apart are recommended for

Total Cholesterol, Triglycerides, HDL & LDL Cholesterol. 2. As per NLA-2014 guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended.

 Jow HDL levels are associated with increased risk for Atherosclerotic Cardiovascular disease (ASCVD) due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated from peripheral tissues.
 NLA-2014 identifies Non HDL Cholesterol (an indicator of all atherogeniclipoproteins such as LDL, VLDL, IDL, Lpa, Chylomicron remnants) along with LDL-cholesterol as co- primary target for cholesterol lowering therapy. Note that major risk factors can modify treatment goals for LDL & Non HDL

5. Additional testing for Apolipoprotein B, hsCRP,Lp(a) & LP-PLA2 should be considered among patients with moderate risk for ASCVD for risk refinement





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| Test Name | | Value | Unit | Biological Reference interva |
| | | | | |
| | | ENDOCRIN YROID STIMULATIN | G HORMONE (TSH) | |
| by CMIA (CHEMILUMIN Frd GENERATION, ULT | ING HORMONE (TSH): SERUN | YROID STIMULATIN A 2.654 | | 0.35 - 5.50 |
| by CMIA (CHEMILUMIN and GENERATION, ULT | ING HORMONE (TSH): SERUN NESCENT MICROPARTICLE IMMUNG RASENSITIVE | YROID STIMULATIN A 2.654 | G HORMONE (TSH) μIU/mL | 0.35 - 5.50 |
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| by CMIA (CHEMILUMIN Brd GENERATION, ULT | ING HORMONE (TSH): SERUN VESCENT MICROPARTICLE IMMUNG RASENSITIVE AGE 0 – 5 DAYS 6 Days – 2 Months | YROID STIMULATIN A 2.654 | G HORMONE (TSH) μIU/mL REFFERENCE RANGE (0.70 – 15.20 0.70 – 11.00 0.70 – 8.40 0.70 – 7.00 | 0.35 - 5.50 |
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| | ING HORMONE (TSH): SERUN VESCENT MICROPARTICLE IMMUNO RASENSITIVE AGE 0 – 5 DAYS 6 Days – 2 Months 3 – 11 Months 1 – 5 Years 6 – 10 Years 11 - 15 > 20 Years (Adults) | YROID STIMULATIN A 2.654 DASSAY) | G HORMONE (TSH) μIU/mL REFFERENCE RANGE (0.70 – 15.20 0.70 – 11.00 0.70 – 8.40 0.70 – 7.00 0.60 – 5.50 0.50 – 5.50 0.27 – 5.50 | 0.35 - 5.50 |

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USE:- TSH controls biosynthesis and release of thyroid harmones T4 & T3. It is a sensitive measure of thyroid function, especially useful in early or subclinical hypothyroidism, before the patient develops any clinical findings or goitre or any other thyroid function abnormality. **INCREASED LEVELS**:

1. Primary or untreated hypothyroidism, may vary from 3 times to more than 100 times normal depending on degree of hypofunction.

2. Hypothyroid patients receiving insufficient thyroid replacement therapy.

3. Hashimotos thyroiditis.

4.DRUGS: Amphetamines, lodine containing agents and dopamine antagonist.

5.Neonatal period, increase in 1st 2-3 days of life due to post-natal surge.

DECREASED LEVELS:

1. Toxic multi-nodular goitre & Thyroiditis.

2. Over replacement of thyroid harmone in treatment of hypothyroidism.

3. Autonomously functioning Thyroid adenoma

4. Secondary pituatary or hypothalmic hypothyroidism

5. Acute psychiatric illness

6.Severe dehydration.



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7.DRUGS: Glucocorticoids, Dopamine, Levodopa, T4 replacement therapy, Anti-thyroid drugs for thyrotoxicosis. 8.Pregnancy: 1st and 2nd Trimester

LIMITATIONS:

1.TSH may be normal in central hypothyroidism, recent rapid correction of hyperthyroidism or hypothyroidism, pregnancy, phenytoin therapy. 2.Autoimmune disorders may produce spurious results.

*** End Of Report **?



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