



M	Dr. Vinay Chopra ID (Pathology & Microbiology) hairman & Consultant Pathologist	Dr. Yugam MD CEO & Consultant	Pathology)	
NAME: Mrs. BEENA IAGE/ GENDER: 41 YRS/FEMAICOLLECTED BY:REFERRED BY:BARCODE NO.: 01512799CLIENT CODE.: KOS DIAGNOSCLIENT ADDRESS: 6349/1, NICH	LE P R R C	ATIENT ID EG. NO./LAB NO. EGISTRATION DATE OLLECTION DATE EPORTING DATE	: 1543030 : 012407090027 : 09/Jul/2024 09:45 AM : 09/Jul/2024 10:25AM : 09/Jul/2024 10:59AM	
Test Name	Value	Unit	Biological Reference interval	
	CLINICAL CHEMIST	RY/BIOCHEMISTR FILE : BASIC	r	
CHOLESTEROL TOTAL: SERUM by CHOLESTEROL OXIDASE PAP	236.15 ^H	mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR = 240.0	
TRIGLYCERIDES: SERUM by GLYCEROL PHOSPHATE OXIDASE (EN	иzymatic) 192.03 ^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	67.37	mg/dL	LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 - 60.0 HIGH HDL: > OR = 60.0	
LDL CHOLESTEROL: SERUM by CALCULATED, SPECTROPHOTOMETR	y 130.37 ^H	mg/dL	OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129.0 BORDERLINE HIGH: 130.0 - 159.0 HIGH: 160.0 - 189.0 VERY HIGH: > OR = 190.0	
NON HDL CHOLESTEROL: SERUM by CALCULATED, SPECTROPHOTOMETR	у 168.78 ^Н	mg/dL	OPTIMAL: < 130.0 ABOVE OPTIMAL: 130.0 - 159.0 BORDERLINE HIGH: 160.0 - 189.0 HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0	
VLDL CHOLESTEROL: SERUM	38.41	mg/dL	0.00 - 45.00	
by CALCULATED, SPECTROPHOTOMETRY TOTAL LIPIDS: SERUM	664.33	mg/dL	350.00 - 700.00	
by CALCULATED, SPECTROPHOTOMETR CHOLESTEROL/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETR	3.51	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0 MODERATE RISK: 7.10 - 11.0 HIGH RISK: > 11.0	
LDL/HDL RATIO: SERUM	1.94	RATIO	LOW RISK: 0.50 - 3.0	
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DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST MBBS , MD (PATHOLOGY)

KOS Central Lab: 6349/1, Nicholson Road, Ambala Cantt -133 001, HaryanaKOS Molecular Lab: IInd Floor, Parry Hotel, Staff Road, Opp. GPO, Ambala Cantt -133 001, Haryana0171-2643898, +91 99910 43898care@koshealthcare.comwww.koshealthcare.com



TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.





	· · · · · ·	hopra & Microbiology) onsultant Patholog		(Pathology)
NAME	: Mrs. BEENA DEVI			
AGE/ GENDER	: 41 YRS/FEMALE		PATIENT ID	: 1543030
COLLECTED BY	:		REG. NO./LAB NO.	: 012407090027
REFERRED BY	:		REGISTRATION DATE	: 09/Jul/2024 09:45 AM
BARCODE NO.	:01512799		COLLECTION DATE	: 09/Jul/2024 10:25AM
CLIENT CODE.	: KOS DIAGNOSTIC LAB		REPORTING DATE	: 09/Jul/2024 10:59AM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD), AMBALA CANT	Т	
Test Name		Value	Unit	Biological Reference interval
by CALCULATED, SPE	ECTROPHOTOMETRY			MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HD by CALCULATED, SP	L RATIO: SERUM ECTROPHOTOMETRY	2.85 ^L	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.

3. Low 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. 4. 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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MBBS, MD (PATHOLOGY)







NAME : Mrs. BEENA DEV AGE/ GENDER : 41 YRS/FEMALE COLLECTED BY : REFERRED BY : BARCODE NO. : 01512799 CLIENT CODE. : KOS DIAGNOSTIC CLIENT ADDRESS : 6349/1, NICHOLS Test Name URIC ACID: SERUM by URICASE - OXIDASE PEROXIDASE INTERPRETATION:- 1.GOUT occurs when high levels of Uric Ac 2.Uric Acid is the end product of purine me intestinal tract by microbial degradation. INCREASED:- (A).DUE TO INCREASED PRODUCTION:- 1.Idiopathic primary gout. 2.Excessive dietary purines (organ meats,le 3.Cytolytic treatment of malignancies espet 4.Polycythemai vera & myeloid metaplasia 5.Psoriasis. 6.Sickle cell anaemia etc. (B).DUE TO DECREASED EXCREATION (BY KIII 1.Alcohol ingestion. 2.Thiazide diuretics. 3.Lactic acidosis. 4.Aspirin ingestion (less than 2 grams per of 5.Diabetic ketoacidosis or starvation. 6.Renal failure due to any cause etc. DECREASED:- (A).DUE TO DIETARY DEFICIENCY 1.Dietary deficiency of Zinc, Iron and molyl 2.Fanconi syndrome & Wilsons disease. 3.Multiple sclerosis . 4.Syndrome of inappropriate antidiuretic h	r. Vinay Chopra D (Pathology & Microbiolog airman & Consultant Patho	y)	Yugam Chopra MD (Pathology) nsultant Pathologist	
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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.





Dr. Vinay ChopraDr. Yugam ChopraMD (Pathology & Microbiology)MD (Pathology)Chairman & Consultant PathologistCEO & Consultant Pathologist						
NAME AGE/ GENDER COLLECTED BY REFERRED BY BARCODE NO. CLIENT CODE. CLIENT ADDRESS	: Mrs. BEENA DEVI : 41 YRS/FEMALE : : : 01512799 : KOS DIAGNOSTIC LAB : 6349/1, NICHOLSON ROAD, AMBALA CANT	PATIENT ID REG. NO./LAB NO. REGISTRATION DATE COLLECTION DATE REPORTING DATE T	: 1543030 : 012407090027 : 09/Jul/2024 09:45 AM : 09/Jul/2024 10:25AM : 09/Jul/2024 12:08PM			
Test Name	Value	Unit	Biological Reference interval			
ENDOCRINOLOGY PROLACTIN PROLACTIN: SERUM 12.008 ng/mL 3 - 25						
by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY) INTERPRETATION: I.Prolactin is secreted by the anterior pituitary gland and controlled by the hypothalamus. 2. The major chemical controlling prolactin is teretion is dopamine, which inhibits prolactin secretion from the pituitary. 3. Physiological function of prolactin is the stimulation of milk production. In normal individuals, the prolactin level rises in response to physiologic stimuli such as sleep, exercise, nipple stimulation, sexual intercourse, hypoglycemia, postpartum period, and also is elevated in the newborn infant. INCREASED (HYPERPROLACTEMIA): 1. Prolactin-secreting pituitary adenoma (prolactinoma, which is 5 times more frequent in females than males). 2. Functional and organic disease of the hypothalamus. 3. Primary hypothyroidism. 4. Section compression of the pituitary stalk. 5. Chest wall lesions and renal failure. 6. Ectopic tumors. 7. DRUGS: Anti-Dopaminergic drugs like antipsychotic drugs, antinausea/antiemetic drugs, Drugs that affect CNS serotonin metabolism, serotonin receptors, or serotonin recuptake (anti-depressants of all classes, ergot derivatives, some illegal drugs such as cannabis). Antihypertensive drugs Opiates, High doses of estrogen or progesterone, anticonvulsants (valporic acid), anti-tuberculous medications (Isoniazid). SIGNIFICANCE: 1. In loss of libido, impotence, infertility, and hypogonadism in males. Postmenopausal and premenopausal women, as well as men, can also suffer from decreased muscle mass and osteporosis. 3. In males, prolactin levels >27 ng/mL in the absence of pregnancy and postpartum lactation are indicative of hyperprolactinemia. 5. Clear symptoms and signs of hyperprolactinemia are often absent in patients with serum prolactin levels >27 ng/mL in the absence of pregnancy and postpartum lactation are indicative of hyperprolactinemia. 5. Clear symptoms and signs of hyperprolactinemia are often absent in patients with serum prolactin levels >270 ng/mL are indicative of the patients with serum prolactin						
evaluated if signs and symptoms of hyperprolactinemia are absent, or pituitary imaging studies are not informative. '						
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