

Jeevan Jyoti HLM

Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

PATHKIND REFERENCE LAB PATHKIND DIAGNOSTICS PVT. LTD.

Plot No. 55-56, Udyog Vihar, Phase IV, Sector-18, Gurugram-122015 E-Mail: care@pathkindlabs.com | Website: www.pathkindlabs.com Customer Care: 75000 75111

1212069325

Processed By

Pathkind Diagnostics Pvt. Ltd.

Barcode No.

162, Lowther Road, Bai Ka Bagh, Prayagraj

Uttar Pradesh-211003

: Ms. ARADHANA SINGH REG-318023 OPD Name Billing Date 04/04/202308:23:36 Age : 36 Yrs Sample Collected on 04/04/2023 12:26:07 Sex : Female Sample Received on 04/04/2023 15:14:06 : P1212100010920 P. ID No. Report Released on 04/04/2023 15:30:05

: 1212230126 **Accession No**

Referring Doctor: SELF

Referred By Ref no.

Report Status - Preliminary Report			
Test Name	Result	Biological Ref. Interval	Unit
	<u>HAEMATOLO</u>	<u>DGY</u>	
Complete Blood Count (CBC)			
Haemoglobin (Hb) Sample: Whole Blood EDTA Method: Photometric measurement	13.3	12.0 - 15.0	gm/dL
Total WBC Count / TLC Sample: Whole Blood EDTA Method: Impedance	4.9	4.0 - 10.0	thou/μL
RBC Count Sample: Whole Blood EDTA Method: Impedance	4.1	3.8 - 4.8	million/μL
PCV / Hematocrit Sample: Whole Blood EDTA Method: Impedance	39.2	36.0 - 46.0	%
MCV Sample: Whole Blood EDTA Method: Calculated	95.0	83.0 - 101.0	fL
MCH Sample: Whole Blood EDTA Method: Calculated	32.2 H	27.0 - 32.0	pg
MCHC Sample: Whole Blood EDTA Method: Calculated	33.9	31.5 - 34.5	g/dL
RDW (Red Cell Distribution Width) Sample: Whole Blood EDTA Method: Calculated	12.1	11.9 - 15.5	%
<u>DLC (Differential Leucocyte Count)</u> Method: Flowcytometry/Microscopy			
Neutrophils Sample: Whole Blood EDTA Method: VCS Technology & Microscopy	61	40 - 80	%
Lymphocytes Sample: Whole Blood EDTA	32	20 - 40	%













Method: VCS Technology & Microscopy



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04/04/202308:23:36 04/04/2023 12:26:07

Age : 36 Yrs Sex : Female

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Test Name	Result	Biological Ref. Interval	Unit
Eosinophils Sample: Whole Blood EDTA Method: VCS Technology & Microscopy	03	01 - 06	%
Monocytes Sample: Whole Blood EDTA Method: VCS Technology & Microscopy	04	02 - 10	%
Basophils Sample: Whole Blood EDTA Method: VCS Technology & Microscopy	00	00 - 02	%
Absolute Neutrophil Count Sample: Whole Blood EDTA	2989	2000 - 7000	/µL
Absolute Lymphocyte Count Sample: Whole Blood EDTA	1568	1000 - 3000	/µL
Absolute Eosinophil Count Sample: Whole Blood EDTA	147	20 - 500	/µL
Absolute Monocyte Count Sample: Whole Blood EDTA	196 L	200 - 1000	/µL
Absolute Basophil Count Sample: Whole Blood EDTA	00 L	20 - 100	/µL
DLC Performed By Sample: Whole Blood EDTA	EDTA Smear		
Platelet Count Sample: Whole Blood EDTA Method: Impedance	209	150 - 410	thou/μL
MPV (Mean Platelet Volume) Sample: Whole Blood EDTA Method: Calculated	10.7	6.8 - 10.9	fL
Sample: Whole Blood EDTA Erythrocyte Sedimentation Rate (ESR) Sample: Whole Blood EDTA	14 H	<12	mm 1st Hour

Sample: Whole Blood EDTA

Method: Modified Westergren Method









Sex

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%

mg/dl

mg/dl

μg/dL

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Test Name Result Biological Ref. Interval Unit

Blood Group

Blood Grouping

"B"

5.2

102.5

Sample: Whole Blood EDTA

Rh (D) Typing Sample: Whole Blood EDTA **POSITIVE**

BIOCHEMISTRY

HbA1C (Glycosylated Hemoglobin)

HbA1c

Sample: Whole Blood EDTA

Method: Turbidimetric inhibition immunoassay

Non Diabetic: < 5.7 %

74 - 106

70 - 140

5.10 - 14.10

Prediabetic Range: 5.7 - 6.4 %

Diabetic Range : >= 6.5 %

Goal of Therapy :<7.0 % Action suggested:>8.0%

<116.0 mq/dL

Mean Plasma Glucose Sample: Whole Blood EDTA

Method: Calculated

102 Fasting Plasma Glucose

Sample: Fluoride Plasma - F

77 Glucose Post-Prandial

Sample: Fluoride Plasma - PP

Method: Hexokinase

Thyroid Profile Total

0.90 Total T3 (Triiodothyronine) 0.80 - 2.00ng/mL

Sample: Serum

Method: ECLIA

Total T4 (Thyroxine) 8.47

Sample: Serum Method: ECLIA

TSH 3rd Generation 2.490 0.270 - 4.200µIU/mL

Sample: Serum

Method: ECLIA







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CYTOLOGY

GYNAECYTOPATHOLOGY REPORT

Gynaec Cyto no. 0032 / 2023

Clinical details : Routine screening No of slides received / prepared : 02 (labelled)

: Conventional pap smear Specimen type Reporting mode : Bethesda system

: Satisfactory without endocervical cells. Specimen adequacy

Descriptive interpretation

1. Non Neoplastic findings : Normal morphology of benign squamous epithelial cells seen with

predominance of superficial and intermediate cells Inflammatory cells

mostly neutrophils and occasional histiocytes seen.

: Normal flora 2. Organisms

: Negative for intraepithelial lesion / Malignancy 3. Epithelial abnormalities

: Negative for intraepithelial lesion or malignancy **Impression**

Disclaimer: Gynaecologial cytology is a screening test that aids in the detection of cervical cancer and cancer precurssor. Both false positive and false negative result can occur. The test should be used at regular intervals, and positive result should be confirmed before definitive therapy.













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Nament Ctatus - Dualinainam Dana

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est Name	Result	Biological Ref. Interval	Unit
ipid Profile			
Total Cholesterol Sample: Serum Method: Spectrophotometery	148	No risk : < 200 Moderate risk : 200–239 High risk : =240	mg/dL
Triglycerides Sample: Serum Method: Spectrophotometry	52	Desirable : < 150 Borderline High : 150 - 199 High : 200 - 499 Very High : >/= 500	mg/dL
LDL Cholesterol (Calculated) Sample: Serum Method: Calculated	96	Optimal : <100 Near Optimal : 100 - 129 Borderline High : 130 - 160 High : 161 - 189 Very High : >/=190	mg/dL
HDL Cholesterol Sample: Serum Method: Spectrophometry	42	Low : < 40 Optimal : 40 - 60 High : > 60	mg/dl
Non HDL Cholesterol Sample: Serum	106	< 130 mg/d	
VLDL Cholesterol Sample: Serum Method: Calculated	10.4	Desirable 10 - 35	mg/dL
Total Cholesterol / HDL Ratio Sample: Serum Method: Calculated	3.52	Low Risk : 3.3 - 4.4 Average Risk : 4.5 - 7.0 Moderate Risk : 7.1 - 11.0 High Risk : > 11.0	
LDL / HDL Ratio Sample: Serum Method: Calculated	2.3	0.5 - 3.0	
(idnov Profilo (KET)		Low Risk : 0.5 - 3.0 Moderate Risk : 3.1 - 6.0 High Risk : > 6.0	

Kidney Profile (KFT)

Blood Urea















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Preliminary Report Donort Status

Report Status - Preliminary Report Test Name Result Biological Ref. Interval Unit				
Blood Urea Nitrogen (BUN) Sample: Serum Method: Spectrophotometry-Urease / GLDH	8.21	7.00 - 18.69	mg/dL	
Urea Sample: Serum Method: Spectrophotometery	17.57	17.00 - 43.00	mg/dL	
Creatinine Sample: Serum Method: Spectrophotometry	0.60	0.50 - 1.10	mg/dL	
BUN Creatinine Ratio Sample: Serum Method: Calculated	14	10 - 20		
Calcium Sample: Serum Method: Spectrophotometery	9.6	8.6 - 10.0	mg/dL	
Uric Acid Sample: Serum Method: Spectrophotometery	3.7	2.4 - 5.7	mg/dL	
Total Protein Sample: Serum Method: Spectrophotometry	7.3	6.4 - 8.3	g/dL	
Albumin Sample: Serum Method: Spectrophotometery	5.1 H	4.0 - 4.9	g/dL	
Globulin Sample: Serum Method: Calculated	2.2	1.9 - 3.7	g/dL	
Albumin/Globulin (A/G) Ratio Sample: Serum	2.3 H	1.0 - 2.1	g/dL	

Method: Calculated









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Report Status - Preliminary Report

Test Name Result Biological Ref. Interval Unit

CLINICAL PATHOLOGY

Urine Routine & Microscopic Examination

Method: Reflectance Photometry

Physical Examination

Colour

Sample: Urine

Method: Physical Examination

Appearance

Sample: Urine

Method: Physical Examination

Specific Gravity

Sample: Urine Method: pKa change of pretreated polyelectrolytes

pН

Sample: Urine

. Method: Double indicator principle

Pale Yellow

Slightly Hazy

1.005

6.5

1.003 - 1.035

Pale Yellow

Clear

Chemical Examination

Glucose

Sample: Urine

. Method: Glucose oxidase/peroxidase

Protein

Sample: Urine

Method: Protein-error-of-indicators principle

Ketones Sample: Urine

Method: Sodium nitroprusside reaction

Sample: Urine Method: Peroxidase

Bilirubin

Sample: Urine Method: Diazo reaction Not Detected

Not Detected

Not Detected

Not Detected

Not Detected

Not Detected

4.7 - 7.5

Not Detected

Not Detected

Not Detected

Not Detected













Age

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Report Status - Preliminary Report

Test Name	Result	Biological Ref. Interval	Unit
Urobilinogen Sample: Urine Method: Ehrlich's reaction	Normal	Normal	
Nitrite Sample: Urine Method: Nitrite Test	Not Detected	Not Detected	
Microscopic Examination Method: Microscopy			
Pus Cells Sample: Urine	2 - 3	0 - 5	/hpf
RBC Sample: Urine	Not Detected	Not Detected	/hpf
Epithelial Cells Sample: Urine	3 - 5	0 - 5	/hpf
Casts Sample: Urine	Not Detected	Not Detected	/hpf
Crystals Sample: Urine	Not Detected	Not Detected	/hpf
Bacteria Sample: Urine	Not Detected	Not Detected	/hpf
Remarks			

Remarks: Microscopic Examination is performed on urine sediment

BIOCHEMISTRY

Electrolytes (Na/K/CI)

Sodium 140 136 - 145

Sample: Serum

Method: ISE

Sample: Urine





mmol/L











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Test Name	Result	Biological Ref. Interval	Unit
Potassium Sample: Serum Method: ISE	4.3	3.5 - 5.1	mmol/L
Chloride Sample: Serum Method: ISE	110 H	97 - 107	mmol/L

Complete Blood Count (CBC)

Clinical Significance:

CBC comprises of estimation of the cellular componenets of blood including RBCs, WBCs and Platelets. Mean corpuscular volume (MCV) is a measure of the size of the average RBC, MCH is a measure of the hemoglobin cointent of the average RBC and MCHC is the hemoglobin concentration per RBC. The red cell distribution width (RDW) is a measure of the degree of variation in RBC size (anisocytosis) and is helpful in distinguishing between some anemias. CBC examination is used as a screening tool to confirm a hematologic disorder, to establish or rule out a diagnosis, to detect an unsuspected hematologic disorder, or to monitor effects of radiation or chemotherapy. Abnormal results may be due to a primary disorder of the cell-producing organs or an underlying disease. Results should be interpreted in conjunction with the patient's clinical picture and appropriate additional testing performed.

Erythrocyte Sedimentation Rate (ESR)

The erythrocyte sedimentation rate (ESR) is a simple but non-specific test that helps to detect inflammation associated with conditions such as infections, cancers, and autoimmune diseases.

HbA1C (Glycosylated Hemoglobin)

Clinical Significance:

Hemoglobin A1c (HbA1c) level reflects the mean glucose concentration over the previous period (approximately 8-12 weeks) and provides a much better indication of long-term glycemic control than blood and urinary glucose determinations. American Diabetes Association (ADA) include the use of HbA1c to diagnose diabetes, using a cutpoint of 6.5%. The ADA recommends measurement of HbA1c 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to assess whether a patient's metabolic control has remained continuously within the target range. Falsely low HbA1c results may be seen in conditions that shorten erythrocyte life span, and may











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Test Name	Result	Biological Ref. Interval	Unit	
		3		

not reflect glycemic control in these cases accurately.

Total T3 (Triiodothyronine)

Clinical Significance:

Thyroid hormones, T3 and T4, which are secreted by the thyroid gland, regulate a number of developmental, metabolic, and neural activities throughout the body. The thyroid gland synthesizes 2 hormones - T3 and T4. T3 production in the thyroid gland constitutes approximately 20% of the total circulating T3, 80% being produced by peripheral conversion from T4. T3 is more potent biologically. Total T3 comprises of Free T3 and bound T3. Bound T3 remains bound to carrier proteins like thyroid-binding globulin, prealbumin, and albumin). Only the free forms are metabolically active. In hyperthyroidism, both T4 and T3 levels are usually elevated, but in some rare cases, only T3 elevation is also seen. In hypothyroidism T4 and T3 levels are both low. T3 levels are frequently low in sick or hospitalized euthyroid patients.

Total T4 (Thyroxine)

Clinical Significance:

Total T4 is synthesized in the thyroid gland. About 0.05% of circulating T4 is in the free or biologically active form. The remainder is bound to thyroxine-binding globulin (TBG), prealbumin, and albumin. High levels of T4 (and FT4) causes hyperthroidism and low levels lead to hypothyroidism.

TSH 3rd Generation

Clinical Significance:

TSH levels are elevated in primary hyporthyroidism and low in primary hyperthyroidism. Evaluation of TSH is useful in the differential diagnosis of primary from secondary and tertiary hypothyroidism. In primary hypothyroidism, TSH levels are elevated, while in secondary and tertiary hypothyroidism, TSH levels are low or normal. High TSH level in the presence of normal FT4 is called subclinical hypothyroidism and low TSH with normal FT4 is called subclinical hyperthyroidism. Sick, hospitalized patients may have falsely low or transiently elevated TSH. Significant diurnal variation is also seen in TSH levels.

Guidelines for TSH levels in pregnancy, as per American Thyroid Association, are as follows:

PREGNANCY TRIMESTER	BIOLOGICAL REFERENCE INTERVAL	UNIT
FIRST TRIMESTER	0.100 - 2.500	μIU/mL
SECOND TRIMESTER	0.200 - 3.000	μIU/mL
THIRD TRIMESTER	0.300 - 3.000	uIU/mL

PAP (Papanicolaou) Smear









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10001141110		g	••

Clinical Significance:

Referred By

PAP smear is used for screening for cervical carcinoma and infections of the female genital tract including human papillomavirus, herpes, Candida, and Trichomonas. Standard reporting done as defined by the Bethesda System (TBS).

Lipid Profile

Proposed LDL-C goals in very high risk and extreme risk group patients by the Lipid Association of India.

Very High Risk group(VHRG)	Extreme Risk group		
	Category A	Category B	
LDL-C goal of <50 mg/dl	LDL-C goal of <50 mg/dl (recommended) LDL-C goal of ≤30 mg/dl (optional)	LDL-C goal of ≤30 mg/dl	
High-risk conditions		CAD with ≥ 1 of following:	
Any one of following:			
 ASCVD (CAD/PAD/TIA or stroke) Homozygous familial hypercholesterolemia Diabetes with ≥2 major ASCVD risk factors*/target organ damage 	CAD with ≥1 of following: 1. Diabetes without target organ damage/≤1 major 2. ASCVD risk factors 3. Familial hypercholesterolemia 4. ≥3 major ASCVD risk factors 5. CKD stage 3B and 4 6. ≥2 major ASCVD risk factors with ≥1 moderate 7. non-conventional risk factor# 8. Lp(a) ≥50 mg/dl 9. Coronary calcium score ≥300 HU 10. Extreme of a single risk factor 11. PAD 12. H/o TIA or stroke 13. Non-stenotic carotid plaque	 Diabetes + polyvascular disease/≥2 major ASCVD risk factors*/target organ damage Recurrent ACS (within 12 months) despite on LDL-C goal Homozygous familial Hypercholesterolemia 	









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Report Status -**Preliminary Report**

Biological Ref. Interval Test Name Result Unit

The LDL-C goal of ≤30 mg/dl must be pursued after detailed risk-benefit discussion between physician and patient.

Clinical judgment to be used in decision making if the patient has disease/risk factors not covered in the table, eg. peripheral arterial disease or cerebrovascular disease.

*Major ASCVD risk factors: 1. Age- male ≥45 years, female ≥55 years, 2. Family h/o premature CAD- male <55 years, female <65 years, 3. Smoking/tobacco use, 4. Systemic hypertension, 5.Low HDL (males <40 mg/dl and females <50 mg/dl).

#Moderate non-conventional risk factors: 1. Coronary calcium score 100–299 HU, 2. Increased carotid intima-media thickness, 3. Lp(a) ≥20–49

Uric Acid

Clinical Significance:

Uric acid is the final product of purine metabolism. Serum uric acid levels are raised in case of increased purine synthesis, inherited metabolic disorder, excess dietary purine intake, increased nucleic acid turnover, malignancy and cytotoxic drugs. Decreased levels are seen in chronic renal failure, severe hepatocellular disease with reduced purine synthesis, defective renal tubular reabsorption, overtreatment of hyperuricemia with allopurinol, as well as some cancer therapies.

Urine Routine & Microscopic Examination

Clinical Significance:

Urine routine examination and microscopy comprises of a set of screening tests that can detect some common diseases like urinary tract infections, kidney disorders, liver problems, diabetes or other metabolic conditions. Physical characteristics (colour and appearance), chemical composition (glucose, protein, ketone, blood, bilirubin and urobilinogen) and microscopic content (pus cells, epithelial cells, RBCs, casts and crystals) are analyzed and reported.

** End of Report**

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