

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

**Processed By** 

### Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

Uttar Pradesh-211003

: Mr. NILESH KUMAR DUBEY REG-309782 OPD **Billing Date** 10/09/202211:15:17 Name Age : 33 Yrs Sample Collected on 11/09/2022 14:28:06

Sex : Male Sample Received on 11/09/2022 17:13:30

P. ID No. : P1212100000371 Report Released on 11/09/2022 17:15:00

: 121222020378 **Accession No** Barcode No. 15457257 Referring Doctor: DEVENDRA TRIPATH

Referred By Ref no.

#### Report Status - Final

Report Status - Final			
Test Name	Result	Biological Ref. Interval	Unit
	HAEMATOLO	<u>DGY</u>	
Complete Blood Count (CBC)			
Haemoglobin (Hb) Sample: Whole Blood EDTA Method: Photometric measurement	15.3	13.0 - 17.0	gm/dL
Total WBC Count / TLC Sample: Whole Blood EDTA Method: Impedance	5.8	4.0 - 10.0	thou/μL
RBC Count Sample: Whole Blood EDTA Method: Impedance	5.2	4.5 - 5.5	million/μL
PCV / Hematocrit Sample: Whole Blood EDTA Method: Impedance	46.6	40.0 - 50.0	%
MCV Sample: Whole Blood EDTA Method: Calculated	88.9	83.0 - 101.0	fL
MCH Sample: Whole Blood EDTA Method: Calculated	29.2	27.0 - 32.0	pg
MCHC Sample: Whole Blood EDTA Method: Calculated	32.8	31.5 - 34.5	g/dL
RDW (Red Cell Distribution Width) Sample: Whole Blood EDTA Method: Calculated	12.2	11.8 - 15.6	%
DLC (Differential Leucocyte Count)  Method: Flowcytometry/Microscopy			
Neutrophils Sample: Whole Blood EDTA Method: VCS Technology & Microscopy	51	40 - 80	%
Lymphocytes Sample: Whole Blood EDTA	42 H	20 - 40	%



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Method: VCS Technology & Microscopy









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Sample Collected on

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Uttar Pradesh-211003

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est 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	2958	2000 - 7000	/μL
Absolute Lymphocyte Count Sample: Whole Blood EDTA	2436	1000 - 3000	/μL
Absolute Eosinophil Count Sample: Whole Blood EDTA	174	20 - 500	/μL
Absolute Monocyte Count Sample: Whole Blood EDTA	232	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	199	150 - 410	thou/μL
MPV (Mean Platelet Volume) Sample: Whole Blood EDTA Method: Calculated	12.9 H	6.8 - 10.9	fL
Sample: Whole Blood EDTA  Tythrocyte Sedimentation Rate (ESR)	14 H	<10	mm 1st Hour

Sample: Whole Blood EDTA

Method: Modified Westergren Method







जांच सही तो इलाज सही (जांच)



P. ID No.

#### Jeevan Jyoti HLM

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15457256, 15457192, 15457255, 15457257

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Test Name	Result	Biological Ref. Interval	Unit
10001101110			•

**Blood Group** 

**Blood Grouping** 

Sample: Whole Blood EDTA

Rh (D) Typing

Sample: Whole Blood EDTA

" O "

Positive

#### **BIOCHEMISTRY**

## **HbA1C (Glycosylated Hemoglobin)**

Method: Turbidimetric inhibition immunoassay

HbA1c	5.6	Non Diabetic : < 5.7 %	%
Sample: Whole Blood EDTA		Prediabetic Range : 5.7 - 6.4 %	

Diabetic Range: >= 6.5 % Goal of Therapy :<7.0 % Action suggested :>8.0 %

70 - 140

5.10 - 14.10

Mean Plasma Glucose 114 <116.0 mg/dL

Sample: Whole Blood EDTA Method: Calculated

**Fasting Plasma Glucose** 89 74 - 106 mg/dl

102

Sample: Fluoride Plasma - F **Glucose Post-Prandial** 

Sample: Fluoride Plasma - PP Method: Hexokinase

#### **Thyroid Profile Total**

Total T3 (Triiodothyronine)	1 21	0.80 - 2.00	ng/mL
iotal 13 (Trilogothyronine)	1.21	0.80 - 2.00	US/UIL

Sample: Serum Method: ECLIA

**Total T4 (Thyroxine)** 10.09

Sample: Serum Method: ECLIA

**TSH 3rd Generation** 2.830 0.270 - 4.200 μIU/mL

Sample: Serum Method: ECLIA



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mg/dl

 $\mu g/dL$ 





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### **Report Status - Final**

Unit Test Name Result **Biological Ref. Interval** 

### **CLINICAL PATHOLOGY**

## **Stool Routine & Microscopic Examination**

# **Physical Examination**

**Brownish** Yellowish Brown Colour

Sample: Stool

**Consistency** Semi Solid Semi Solid

Sample: Stool

Absent Absent Mucus

Sample: Stool

Absent Absent Blood

Sample: Stool

Odour **Fecal Fecal** Sample: Stool

#### **Microscopic Examination**

Not Detected Not Detected Cyst

Not Detected **Trophozoites** Not Detected

Sample: Stool

Sample: Stool

**Charcot - Leyden Crystals** Not Detected Not Detected

Sample: Stool

Sample: Stool

Not Detected Not Detected Ova

Not Detected Not Detected **Adult Parasite** 

Sample: Stool



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Age

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: 33 Yrs

Sex : Male P. ID No. : P1212100000371

Sample: Stool

Accession No : 121222020378

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Test Name	Result	Biological Ref. Interval	Unit
RBC Sample: Stool	Not Detected	0 - 0	/hpf
Pus Cells Sample: Stool	1 - 2	0 - 5	/HPF
Stool pH & Reducing Substances			
Stool for pH Sample: Stool	6.5		
Stool For Reducing Substances	Not Detected	Not Detected	











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Test Name	Result	Biological Ref. Interval	Unit
	BIOCHEMIS <sup>*</sup>	<u>TRY</u>	
<b>Liver Function Test (LFT)</b>			
Bilirubin Total Sample: Serum Method: Spectrophotometery	0.6	<1.1	mg/dL
Bilirubin Direct Sample: Serum Method: Spectrophotometery	0.2	<0.2	mg/dL
<b>Serum Bilirubin (Indirect)</b> Sample: Serum Method: Calculated	0.40	<0.90	mg/dL
SGOT / AST Sample: Serum Method: Spectrophotometery	35	<37	U/L
SGPT / ALT Sample: Serum Method: Spectrophotometery	55 H	<41	U/L
AST / ALT Ratio Sample: Serum Method: Calculated	0.64		
Alkaline Phosphatase (ALP) Sample: Serum Method: Spectrophotometery	141 H	<128	U/L
<b>Total Protein</b> Sample: Serum Method: Spectrophotometry	8.9 H	6.4 - 8.3	g/dL
<b>Albumin</b> Sample: Serum Method: Spectrophotometery	5.4 H	4.0 - 4.9	g/dL
<b>Globulin</b> Sample: Serum Method: Calculated	3.5	1.9 - 3.7	g/dL



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Report Status - Final			
Test Name	Result	Biological Ref. Interval	Unit
Albumin/Globulin (A/G) Ratio Sample: Serum Method: Calculated	1.5	1.0 - 2.1	g/dL
Lipid Profile			
<b>Total Cholesterol</b> Sample: Serum Method: Spectrophotometery	198	No risk : < 200 Moderate risk : 200–239 High risk : =240	mg/dL
<b>Triglycerides</b> Sample: Serum Method: Spectrophotometry	111	Desirable : < 150 Borderline High : 150 - 199 High : 200 - 499 Very High : >/= 500	mg/dL
LDL Cholesterol (Calculated) Sample: Serum Method: Calculated	137 H	Optimal : <100 Near Optimal : 100 - 129 Borderline High : 130 - 160 High : 161 - 189 Very High : >/=190	mg/dL
HDL Cholesterol Sample: Serum Method: Spectrophometry	39 L	Low : < 40 Optimal : 40 - 60 High : > 60	mg/dl
Non HDL Cholesterol Sample: Serum	159 H	< 130	mg/dL
VLDL Cholesterol Sample: Serum Method: Calculated	22.2	Desirable 10 - 35	mg/dL
<b>Total Cholesterol / HDL Ratio</b> Sample: Serum Method: Calculated	5.08 H	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	3.5 H	0.5 - 3.0	

Low Risk : 0.5 - 3.0



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Method: Calculated



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Report Status - Tillai			
Test Name	Result	Biological Ref. Interval	Unit
Kidney Profile (KFT) Blood Urea		Moderate Risk : 3.1 - 6.0 High Risk : > 6.0	
Blood Urea Nitrogen (BUN) Sample: Serum Method: Spectrophotometry-Urease / GLDH	8.58 L	8.87 - 20.50	mg/dL
<b>Urea</b> Sample: Serum Method: Spectrophotometery	18.36	17.00 - 43.00	mg/dL
Creatinine Sample: Serum Method: Spectrophotometry	0.97	0.70 - 1.30	mg/dL
BUN Creatinine Ratio Sample: Serum Method: Calculated	9 L	10 - 20	
<b>Calcium</b> Sample: Serum Method: Spectrophotometery	10.2 H	8.6 - 10.0	mg/dL
<b>Uric Acid</b> Sample: Serum Method: Spectrophotometery	8.8 H	3.4 - 7.0	mg/dL
<b>Total Protein</b> Sample: Serum Method: Spectrophotometry	8.9 H	6.4 - 8.3	g/dL
<b>Albumin</b> Sample: Serum Method: Spectrophotometery	5.4 H	4.0 - 4.9	g/dL
<b>Globulin</b> Sample: Serum Method: Calculated	3.5	1.9 - 3.7	g/dL
Albumin/Globulin (A/G) Ratio Sample: Serum	1.5	1.0 - 2.1	g/dL







Method: Calculated







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Pale Yellow

**Report Status - Final** 

Test Name Result **Biological Ref. Interval** Unit

**CLINICAL PATHOLOGY** 

**Urine Routine & Microscopic Examination** 

Method: Reflectance Photometry

**Physical Examination** 

Pale Yellow Colour

Sample: Urine

Method: Physical Examination

Clear Clear **Appearance** 

Sample: Urine

Method: Physical Examination

1.010 Specific Gravity 1.003 - 1.035

Sample: Urine

Method: pKa change of pretreated polyelectrolytes

6.0 4.7 - 7.5 pН

Sample: Urine

Method: Double indicator principle

**Chemical Examination** 

Not Detected Not Detected Glucose

Sample: Urine

Method: Glucose oxidase/peroxidase

**Protein** Not Detected Not Detected

Sample: Urine

Method: Protein-error-of-indicators principle

Not Detected Not Detected Ketones

Sample: Urine

Method: Sodium nitroprusside reaction

Not Detected Not Detected Blood

Sample: Urine

Method: Peroxidase

Bilirubin Not Detected Not Detected

Sample: Urine Method: Diazo reaction



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Test Name	Result	Biological Ref. Interval	Unit
<b>Urobilinogen</b> Sample: Urine Method: Ehrlich's reaction	Normal	Normal	
<b>Nitrite</b> Sample: Urine Method: Nitrite Test	Not Detected	Not Detected	
Microscopic Examination  Method: Microscopy			
Pus Cells Sample: Urine	1 - 2	0 - 5	/hpf
RBC Sample: Urine	Not Detected	Not Detected	/hpf
<b>Epithelial Cells</b> Sample: Urine	1 - 2	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 141 136 - 145 mmol/L

Sample: Serum Method: ISE

Sample: Urine













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<b>Accession No</b>	: 121222020378	Barcode No.	:	15457256, 15457192,
Referring Docto	: DEVENDRA TRIPATH			15457255, 15457258, 15457259, 15457257
Referred By	:	Ref no.	:	15457255, 15457257

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est Name	Result	Biological Ref. Interval	Unit
Potassium Sample: Serum Method: ISE	4.4	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
		<del>-</del>	

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.

### **Stool Routine & Microscopic Examination**

### Clinical Significance:

Routine and microscopic examination of stool sample comprises of macroscopic as well as microscopic examination of the sample for presence of parasitic ova and cysts.

#### Stool for pH













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# Pathkind Diagnostics Pvt. Ltd.

162, Lowther Road, Bai Ka Bagh, Prayagraj

Uttar Pradesh-211003

Ref no.

: Mr. NILESH KUMAR DUBEY REG-309782 OPD **Billing Date** 10/09/202211:15:17 Name : 33 Yrs Sample Collected on 11/09/2022 14:28:06 Age Sex Male Sample Received on 11/09/2022 17:13:30 P. ID No. : P1212100000371 Report Released on 11/09/2022 17:15:00 **Accession No** : 121222020378 Barcode No. 15457256, 15457192, 15457255, 15457258, Referring Doctor: DEVENDRA TRIPATH

15457259, 15457257

**Report Status - Final** 

Test Name Result **Biological Ref. Interval** Unit

#### Clinical Significance:

Testing for pH and reducing substances in stool helps in determining the underlying cause of diarrhea - whether the diarrhoea is due to osmotic cause or due to infective cause.

#### **Bilirubin Total**

"Total Bilirubin is one of the most commonly used tests to assess liver function. A number of inherited and acquired diseases affect bilirubin production, metabolism, storage and excretion and causes hyperbilirubinemia resulting in jaundice. Hyperbilirubinemia may be due to increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Unconjugated hyperbilirubinemia is seen in newborn andd known as physiological jaundice. Elevated unconjugated bilirubin in the neonatal period may result in brain damage (kernicterus). Crigler-Najjar syndromes type I and type II are also associated with elevated levels of indirect bilirubin. Both conjugated and unconjugated bilirubin are increased in hepatitis and space-occupying lesions of the liver; and obstructive lesions such as carcinoma of the head of the pancreas, common bile duct, or ampulla of Vater."

#### **Bilirubin Direct**

### Clinical Significance:

"Direct bilirubin is a measurement of conjugated bilirubin. Jaundice can occur as a result of increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Inherited disorders in which direct bilirubin levels are increased are seen in Dubin-Johnson syndrome and Rotor syndrome, idiopathic neonatal hepatitis and biliary atresia. The most commonly occurring form of jaundice of the newborn called physiological jaundiceis due to increase in levels of indirect bilirubin. Both conjugated and unconjugated bilirubin are increased in hepatocellular diseases such as hepatitis and space-occupying lesions of the liver, bstructive lesions such as carcinoma of the head of the pancreas, common bile duct, or ampulla of Vater."

#### SGOT / AST

#### Clinical Significance:

"Elevated aspartate aminotransferase (AST) values are seen most commonly in parenchymal liver diseases. Values can be elevated from 10 to 100 times the normal range, though commonly 20 to 50 times elevations are seen. AST levels are raised in infectious hepatitis and other inflammatory conditions affecting the liver along with ALT, though ALT levels are higher. The ALT:AST ratio which is normally <1 is reversed in these conditions











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and becomes >1. AST levels are usually raised before clinical signs and symptoms of disease appear. AST and ALT also rise in primary or metastatic carcinoma of the liver, with AST usually being higher than ALT. Elevated AST values may also be seen in disorders affecting the heart, skeletal muscle and kidney, such as myocardial infarction, muscular dystrophy, dermatomyositis, acute pancreatitis and crushed muscle injuries."

#### SGPT / ALT

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#### Clinical Significance:

Elevated alanine aminotransferase (ALT) values are seen in parenchymal liver diseases characterized by a destruction of hepatocytes. Values are at least 10 times higher the normal range and may reach up to 100 times the upper reference limit. Commonly, values are seen to be 20 - 50 times higher than normal. In infectious hepatitis and other inflammatory conditions affecting the liver, ALT levels rise more than aspartate aminotransferase (AST), and the ALT/AST ratio, which is normally <1, is reversed and becomes >1. ALT levels usually rise before clinical signs and symptoms of disease appear.

#### Alkaline Phosphatase (ALP)

#### Clinical Significance:

Alkaline Phosphatase levels can be elevated in both liver related as well as bone related conditions. ALP levels are raised (more than 3 fold) in extrahepatic biliary obstruction (eg, by stone or by cancer of the head of the pancreas) than in intrahepatic obstruction, and is directly proportional to the level of obstruction. Levels may rise up to 10 to 12 times the upper limit of normal range and returns to normal on surgical removal of the obstruction. ALP levels rise together with GGT levels and If both GGT and ALP are elevated, a liver source of the ALP is likely. Among bone diseases, ALP levels rise in Paget disease (up to 25 fold), osteomalacia, rickets, primary and secondary hyperparathyroidism and osteogenic bone cancer. Elevated ALP is seen in children following accelerated bone growth. Also, a 2 to 3fold elevation may be observed in women in the third trimester of pregnancy, although the interval is very wide and levels may not exceed the upper limit of the reference interval in some cases.

# **Total Protein**

### Clinical Significance:

High levels of Serum Total Protein is seen in increased acute phase reactants in inflammation, late-stage liver disease, infections, multiple myeloma and other malignant paraproteinemias.n. Hypoproteinemia is seen in hypogammaglobulinemia, nephrotic syndrome and protein-losing enteropathy.

#### **Albumin**









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Referring Docto	or : DEVENDRA TRIPATH	15457255, 15457258, 15457259, 15457257
Referred By	:	Ref no. :

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### Clinical Significance:

"Hypoalbuminemia can be caused by impaired synthesis due to liver disease (primary) or due to diminished protein intake (secondary), increased catabolism due to tissue damage and inflammation; malabsorption of amino acids; and increased renal excretion (eg, nephrotic syndrome). Hyperalbuminemia is seen in dehydration."

# **Lipid Profile**

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

Extreme Risk group		
Category A	Category B	
LDL-C goal of <50 mg/dl (recommended) LDL-C goal of ≤30 mg/dl (optional)	LDL-C goal of ≤30 mg/dl	
	CAD with $\geq 1$ of following:	
CAD with ≥1 of following:	<ol> <li>Diabetes + polyvascular disease/≥2</li> <li>major ASCVD risk factors*/target</li> </ol>	
<ol> <li>Diabetes without target organ damage/≤1 major</li> <li>ASCVD risk factors</li> <li>Familial hypercholesterolemia</li> <li>≥3 major ASCVD risk factors</li> <li>CKD stage 3B and 4</li> <li>≥2 major ASCVD risk factors with ≥1 moderate</li> <li>non-conventional risk factor#</li> <li>Lp(a) ≥50 mg/dl</li> <li>Coronary calcium score ≥300 HU</li> <li>Extreme of a single risk factor</li> </ol>	organ 3. damage 4. Recurrent ACS (within 12 months) 5. despite on LDL-C goal 6. Homozygous familial 7. Hypercholesterolemia	
	Category A  LDL-C goal of <50 mg/dl (recommended)  LDL-C goal of ≤30 mg/dl (optional)  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	









#### Jeevan Jyoti HLM

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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\*\*

Dr. Ankit Singh

MBBS, MD (Pathologist) Lab Head











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DIABETES	FBS, HbA1c	FBS, HbA1c, Microalbumin	FBS, HbA1c, Microalbumin
KIDNEY	BUN, Creatinine, Bun/Creatinine Ratio, Electrolytes, Uric Acid, Urine R/E	BUN, Creatinine, BUN/Creatinine Ratio, Electrolytes, Uric Acid, Urine R/E	BUN, Creatinine, BUN/Creatinine Ratio, Electrolytes, Uric Acid, Urine R/E
BONES	Vitamin D, Calcium	Vitamin D, Calcium, Phosphorus	Vitamin D, Calcium, Phosphorus, Rheumatoid Factor
THYROID	T3, T4, TSH	T3, T4, TSH	FT3, FT4, TSH
NERVES	Vitamin B12	Vitamin B12	Vitamin B12
LIVER	Bilirubin (Total, Direct, Indirect), SGOT, SGPT, ALP, Protein, Albumin, Globulin, A:G Ratio, HBsAg	Bilirubin (Total, Direct, Indirect), SGOT, SGPT, ALP, GGT, LDH, Protein, Albumin, Globulin, A:G Ratio, HBsAg	Bilirubin (Total, Direct, Indirect), SGOT, SGPT, ALP, GGT, LDH, Protein, Albumin, Globulin, A:G Ratio, HBsAg
ANAEMIA	Iron, TIBC, UIBC, % Saturation	Iron, TIBC, UIBC, % Saturation, Ferritin	Iron, TIBC, UIBC, % Saturation, Ferritin, Folic Acid
INFECTION	CBC, ESR	CBC, ESR	CBC, ESR

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