



Name : Mr. MANOJ KUMAR UHID : 117657 S No : PID : 30302  
Age/Gender : 38 Year/Male A.S : NP Sample Date : 11-Jun-2024 10:54 AM  
Ref. By Dr. : MEDIWHEEL Report Date : 11-Jun-2024 05:11 PM  
Address : HISAR Sample Type : Inside \*30302\*

Test Name	Value	Unit	Reference Range
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## HEAMATOLOGY

### CBC (Complete Blood Count)

Haemoglobin (Hb)	13.7	g/dl	12.0 - 17.4 g/dl
<b>Total RBC Count</b>	<b>6.23</b>	m/cumm	4.70 - 6.10
Haematocrit	43.1	%	35.0 - 50.0 %
<b>Mean Cell Volume</b>	<b>69.2</b>	fL	80.0 - 100 fL
<b>Mean Cell Haemoglobin</b>	<b>22.0</b>	pg	27.0 - 34.0 pg
<b>Mean Cell Haemoglobin Conc</b>	<b>31.8</b>	%	32.0 - 36.0
Red Cell Distribution Width (RDW)-CV	14.5	%	11.0 - 16.0 %
Red Cell Distribution Width (RDW)-SD	40.2	fL	35.0 - 56.0 fL
-			
Total Leucocyte Count	5970	cells/cum m	4000 - 11000
Differential Leucocyte Count	.		
Neutrophils	45	%	32 - 72 %
Lymphocytes	50	%	20 - 50 %
Monocytes	3	%	2 - 11 %
Eosinophils	2	%	1 - 3 %
Basophils	0	%	0 - 2 %
Platelet Count	1,99,000	cells/cunm m	150,000 - 450,000
Platelet Distribution Width	15.1	fL	15.0 - 18.0 fL
Mean Platelet Volume	11.2	fL	7.0 - 13.0 fL

Sample Type : Whole Blood

1. Spurious elevation of platelet count may be seen in patients with extensive burns, extreme microcytosis, microangiopathic hemolytic anemia, red cell fragmentation, micro-organisms like bacteria, fungi or yeast, hyperlipidemia, fragments of white blood cell (WBC) cytoplasm in patients with acute leukemia, hairy cell leukemia, lymphomas and in presence of cryoglobulins.
2. Spuriously low platelet counts may be seen in cases of platelet clumping (EDTA induced, platelet cold agglutinins, multiple myeloma), platelet satellitism and in giant platelet syndromes.
3. Delay in processing due to sample transport may cause a mild time dependent fall in platelet count. It is advisable to repeat the test using a citrate / heparin collection tube to avoid this pitfall.
4. Automated platelet counting is subject to 10-15% variation in the result on the same as well as different analysers due to various preanalytic variables like the sampling site, skill in sample collection, anticoagulant used, sample mixing and sample transport etc.

### ABO Blood Grouping

Blood Group

O<sup>+</sup> POSITIVE

Haemaagglutination reaction

A Rh Positive, B Rh Positive, AB Rh Positive, O Rh Positive, A Rh Negative, B Rh Negative, AB Rh Negative, O Rh Negative

Sample Type : Whole Blood

### HBA1C



# Lotus Diagnostic & Imaging Centre

A Unit of Lotus Diagnostic & Imaging Solution Pvt. Ltd.

HB से लेकर MRI तक एक ही छत के नीचे

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Test Name	Value	Unit	Reference Range
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## HBA1C

HBA1C	5.0	%	4.27 - 6.00 %
turbidimetric immunoassay			
Average Blood Glucose	96.8	mg/dl	90.00 - 120.00 mg/dl
turbidimetric immunoassay			
Sample Type : Whole Blood			

### Remarks :

GLYCOSYLATED HEMOGLOBIN (HbA1c)

Reference Range : Please correlate with clinical conditions.

Bellow 6.0 % Normal value

6.0 %-7.0 % Good control

7.0 %-8.0 % Fair control

8.0 %-10 % Unsatisfactory control

Above10 % Poor control

Technology : Immunoassay and chemistry technology to measure A1C and total HB (A1C now Bayer)

AVERAGE BLOOD GLUCOSE (ABG) CALCULATED

Reference Range: Please correlate with clinical conditions.

90-120 mg/dl Excellent control

121-150 mg/d Good control

151-180 mg/dl Average control

181-210 mg/dl Action suggested

> 211 mg/dl Panic values

NOTE: Average blood glucose value is calculated from HbA1C value and it indicates average blood sugar level over past three months.

Technology: Derived from Hb A1C Values

Sample Type: Sodium heparin:

## ESR

ESR	6	mmHr	0 - 15 mmHr
Sample Type : Whole Blood			

Dr. (Maj.)Guruprasad  
MBBS, DMRD, DNB  
Consultant Radiologist

Dr. Rambaksh Sharma  
MBBS, MD  
Consultant Radiologist

Dr. RAJESH REDDU  
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MBBS, MD  
Consultant Pathologist



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Test Name	Value	Unit	Reference Range
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#### CLINICAL COMMENTS:

Erythrocyte sedimentation rate (ESR or sed rate) is a relatively simple, inexpensive, non-specific test that indirectly measures the degree of inflammation present in the body. Inflammation is part of the body's immune response. It can be acute, developing rapidly after trauma, injury or infection, for example, or can occur over an extended time (chronic) with conditions such as autoimmune diseases or cancer.

Moderately elevated ESR occurs with inflammation but also with anemia, infection, pregnancy, and with aging. A very high ESR usually has an obvious cause, such as a severe infection, marked by an increase in globulins, systemic vasculitis, polymyalgia rheumatica or temporal arteritis. People with multiple myeloma or Waldenstrom's macroglobulinemia (tumors that make large amounts of immunoglobulins) typically have very high ESRs even if they don't have inflammation.

#### Factors increasing ESR:

- Advanced age
- Anemia
- Pregnancy
- High fibrinogen
- Macrocytosis
- Kidney problems
- Thyroid disease
- Some cancers, such as multiple myeloma
- Infection

#### Factors decreasing ESR

- Microcytosis
- Low fibrinogen
- Polycythemia
- Marked leukocytosis

### CLINICAL-CHEMISTRY

#### URIC ACID

Uric acid	4.8	mg/dL	3.5 - 7.2
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Uricase - POD  
Sample Type : SERUM

URIC ACID: Increases in case of renal failure, disseminated neoplasms, pregnancy toxemia, psoriasis, liver disease, sarcoidosis etc. Decrease is reported in Wilson's disease, Fanconi's syndrome, xanthinuria.

#### Glucose.Fasting

Glucose, Fasting	87.6	mg/dl	70 - 110 mg/dl
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Hexokinase / GOD - POD

Glucose, Post Prandial	108.3	mg/dl	70 - 140 mg/dl
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Hexokinase / GOD - POD

Sample Type : SERUM



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Test Name	Value	Unit	Reference Range
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Criteria for the diagnosis of diabetes (American diabetes association, 2019)

- Fasting Plasma Glucose  $\geq 126$  mg/dL. Fasting is defined as no caloric intake for at least 8 h.  
OR
- 2-h PG  $\geq 200$  mg/dL during OGTT. The test should be performed using a glucose load containing the equivalent of 75-g anhydrous glucose dissolved in water.\*  
OR
- HbA1c  $\geq 6.5\%$ .  
OR
- Random plasma glucose  $\geq 200$  mg/dL in a patient with classic symptoms of hyperglycemia or hyperglycemic crisis.

Criteria defining prediabetes (American diabetes association, 2019)

- FPG 100 mg/dL to 125 mg/dL (Impaired fasting glucose, IFG)  
OR
- 2-h PG during 75-g OGTT 140 mg/dL to 199 mg/dL (Impaired glucose tolerance, IGT)  
OR
- HbA1c 5.7-6.4%

Note:

All abnormal results must be confirmed with a repeat test on a different day.

## Total Protein

Total Protein	7.1	gm/dl	6.0 - 8.3
BIURET			
Albumin	4.2	g/dl	2.9 - 4.5
BCG			
Globulin	2.9	gm/dl	2.0 - 3.5
Albumin-Globulin Ratio	1.3		1.2 - 2.5

Sample Type : SERUM

## UREA. SERUM

UREA	24.8	mg/dL	14 - 51
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KINETIC METHOD WITH UREASE AND GLDH  
Sample Type : SERUM

UREA: High urea levels suggest poor kidney function, congestive heart failure, shock, stress, recent heart attack or severe burns; bleeding from the gastrointestinal tract; conditions that cause obstruction of urine flow; or dehydration.

Low urea levels can be seen in severe liver disease or malnutrition but are not used to diagnose or monitor these conditions. Low urea levels are also seen in normal pregnancy.

## CREATININE SERUM

CREATININE SERUM	1.0	mg/dL	0.5 - 1.4 mg/dL
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Jaffe Kinetic  
Sample Type : SERUM



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Test Name Value Unit Reference Range

CREATININE: Increases in any renal functional impairment (intrinsic renal lesions, decreased perfusion of the kidney, or obstruction of the lower urinary tract), acromegaly and hyperthyroidism. Decreases in pregnancy, muscle wasting.

**LIVER FUNCTION TEST (LFT) (S)**

Total Bilirubin-Serum	0.90	mg/dl	0.20 - 1.00 mg/dl
Bilirubin Direct Serum	0.40	mg/dl	0.10 - 0.50 mg/dl
Bilirubin Indirect-Serum	0.50	mg/dl	0.20 - 0.70 mg/dl
SGOT	29.4	IU/L	10 - 40 IU/L
IFCC with Pyridoxal Phosphate SGPT	40.7	IU/L	07 - 56 IU/L
IFCC with Pyridoxal Phosphate Alkaline Phosphatase	87.2	U/L	44 - 147 U/L
IFCC PNPP Buffer Total Protein	7.1	gm/dl	6.0 - 8.3
BIURET Albumin	4.2	g/dl	3.5 - 5.5 g/dl
BCG Globulin	2.9	gm/dl	2.0 - 3.5 gm/dl
AG RATIO	1.79		1.2 - 2.5

Sample Type : SERUM

**CLINICAL COMMENT:**

Liver function tests can be suggested in case of hepatitis, liver cirrhosis and monitor possible side effects of medications. A variety of diseases and infections can cause acute or chronic damage to the liver, causing inflammation

(hepatitis), scarring (cirrhosis), bile duct obstructions, liver tumors, and liver dysfunction. Alcohol, drugs, some herbal supplements, and toxins can also injure the liver. A significant amount of liver damage may occur before symptoms such as jaundice, dark urine, light-colored stools, itching (pruritus), nausea, fatigue, diarrhea, and unexplained weight loss or gain appear. Early detection of liver injury is essential in order to minimize damage and preserve liver function.

Alanine aminotransferase (ALT) A very high level of ALT is frequently seen with acute hepatitis. Moderate increases may be seen with chronic hepatitis. People with blocked bile ducts, cirrhosis, and liver cancer may have ALT concentrations that are only moderately elevated or close to normal. Aspartate aminotransferase (AST) A very high level of AST is frequently seen with acute hepatitis. AST may be normal to moderately increased with chronic hepatitis. In people with blocked bile ducts, cirrhosis, and liver cancer, AST concentrations may be moderately increased or close to normal. When liver damage is due to alcohol, AST often increases much more than ALT (this is a

pattern seen with few other liver diseases). AST is also increased after heart attacks and with muscle injury.

AST is a less sensitive and less specific marker of liver injury than ALT. AST is more elevated than ALT in alcohol-induced liver injury. AST could be elevated more than ALT like: (i)

**Lipid Profile**

Cholesterol	162.02	mg/dl	<200.0 mg/dl
CHOD - PAP			
Triglycerides	156.1	mg/dl	< 150 mg/dl
GPO - PAP			
HDL Cholesterol	43.6	mg/dl	Adult males >45 mg/dl
Homoogeneous Enzymatic Colorimetric test			



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Test Name	Value	Unit	Reference Range
<b>Lipid Profile</b>			
LDL Cholesterol	87.2	mg/dl	<100 mg/dl
VLDL Cholesterol	31.22	mg/dl	<30.0 mg/dl
CHO/HDL Ratio	3.72	mg/dl	Low risk 3.3-4.4
Non HDL Cholesterol	118.42	mg/dl	<130 mg/dl

Calculated  
Sample Type : SERUM

#### Interpretation

#### Note

- Measurements in the same patient can show physiological & analytical variations. 3 serial samples 1 wk apart are recommended for Total Cholesterol, Triglycerides, HDL & LDL Cholesterol.
- NLA-2014 identifies Non HDL Cholesterol (an indicator of all atherogenic lipoproteins such as LDL, VLDL, IDL, Lp(a), 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.
- Apolipoprotein B is an optional, secondary lipid target for treatment once LDL & Non HDL goals have been achieved.
- Additional testing for Apolipoprotein B, hsCRP, Lp(a) & LP-PLA2 should be considered among patients with moderate risk for ASCVD for risk refinement.

## CLINICAL PATHOLOGY

### PHYSICAL EXAMINATION

Colour : PALE YELLOW  
Pale-yellow, Yellowish, Colorless, YELLOW  
Quantity : 40 ml  
pH : 6.5  
Mucus : ABSENT  
Absent, Present  
Appearance : CLEAR  
Slightly turbid, Turbid, Clear

### Chemical Examination (Strip)

Specific Gravity : 1.025  
Albumin : NEGATIVE  
Absent, Present(+), Present(2+), Present(3+)  
Sugar : NEGATIVE  
Absent, Present(+), Present(2+), Present(3+)  
Bilirubin : NEGATIVE  
Absent, Present

### Microscopic Examination (Microscopy)

Pus Cells : 2-4 /HPF  
Epithelial Cells : 0-1 /HPF  
RBC : NIL /HPF  
Casts : ABSENT  
Crystals : ABSENT  
Bacteria : ABSENT  
Others :



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Test Name	Value	Unit	Reference Range
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Sample Type : Urine

## ENDOCRINE

### Thyroid Hormones (T3 .T4 & TSH)

T3	0.97	ng/ml	0.60 - 1.81 ng/ml
T4	10.11	ng/dl	5.01 - 12.45 ng/dl
TSH Ultrasensitive	2.12	uIU/ml	0.34 - 5.50 uIU/ml

Sample Type : SERUM

#### Remarks :

Note1. 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 50 %, hence time of the day has influence on the measured serum TSH concentrations.

2. Recommended test for T3 and T4 unbound or free level as it is metabolically active.

3. Physiological rise in Total T3 and T4 level is seen in pregnancy and in patients on steroid therapy.

#### Clinical Use-

- \* Primary Hypothyroidism
- \* Hyperthyroidism
- \* Hypothalamic- Pituitary hypothyroidism
- \* Inappropriate-TSH secretion
- \* Nonthyroidal illness
- \* Autoimmune thyroid disease
- \* Pregnancy associated thyroid disorders
- \* Thyroid dysfunction in infancy and early childhood

--End of Report--