



Name : Mr. JAGDISH KUMAR CHANDA S/o UHID : 111993 S No : PID : 24063
Age/Gender : 56 Year/Male A.S : NP Sample Date : 8-Mar-2024 10:14 AM
Ref. By Dr. : MEDIWHEEL Report Date : 8-Mar-2024 04:52 PM
Address : HISAR Sample Type : Inside *24063*

Test Name	Value	Unit	Reference Range
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HEAMATOLOGY

CBC (Complete Blood Count)

Haemoglobin (Hb)	13.2	g/dl	12.0 - 17.4 g/dl
Total RBC Count	4.55	m/cumm	4.70 - 6.10
Haematocrit	38.1	%	35.0 - 50.0 %
Mean Cell Volume	83.7	fL	80.0 - 100 fL
Mean Cell Haemoglobin	29.0	pg	27.0 - 34.0 pg
Mean Cell Haemoglobin Conc	34.6	%	32.0 - 36.0
Red Cell Distribution Width (RDW) - SD	46.4	fL	35.0 - 56.0 fL
Red Cell Distribution Width (RDW) - CV	12.8	%	11.0 - 16.0 %
Total Leucocyte Count	9680	cells/cum m	4000 - 11000
Differential Leucocyte Count	.		
Neutrophils	70	%	32 - 72 %
Lymphocytes	25	%	20 - 50 %
Monocytes	03	%	2 - 11 %
Eosinophils	02	%	1 - 3 %
Basophils	0	%	0 - 2 %
Platelet Count	2,30,000	cells/cunm m	150,000 - 450,000
Platelet Distribution Width	15.3	fL	15.0 - 18.0 fL
Mean Platelet Volume	11.4	fL	7.0 - 13.0 fL

Sample Type : Whole Blood

- 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.
- 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.
- 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.
- 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

A"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

HBA1C

7.8 % 4.27 - 6.00 %

turbidimetric immunoassay



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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HBA1C

Average Blood Glucose	176.16	mg/dl	90.00 - 120.00 mg/dl
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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	46	mmHr	0 - 15 mmHr
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Sample Type : Whole Blood

Dr. (Maj.)Guruprasad
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Consultant Radiologist

Dr. Rambaksh Sharma
MBBS, MD
Consultant Radiologist

Dr. RAJESH REDDU
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Consultant Physician

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MBBS, MD
Consultant Pathologist



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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	7.6	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, Post Prandial	398.4	mg/dl	70 - 140 mg/dl
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Hexokinase / GOD - POD

Sample Type : SERUM



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Criteria for the diagnosis of diabetes (American diabetes association, 2019)

- Fasting Plasma Glucose ≥ 126 mg/dL. Fasting is defined as no caloric intake for at least 8 h.
OR
- 2-h PG ≥ 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 ≥ 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	6.8	gm/dl	6.0 - 8.3
BIURET			
Albumin	3.98	g/dl	2.9 - 4.5
BCG			
Globulin	2.82	gm/dl	2.0 - 3.5
Albumin-Globulin Ratio	1.16		1.2 - 2.5
Sample Type : SERUM			

CREATININE SERUM

CREATININE SERUM	0.8	mg/dL	0.5 - 1.4 mg/dL
Jaffe Kinetic			
Sample Type : SERUM			

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	54.42	IU/L	10 - 40 IU/L
IFCC with Pyridoxal Phosphate			
SGPT	40.81	IU/L	07 - 56 IU/L
IFCC with Pyridoxal Phosphate			



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Test Name	Value	Unit	Reference Range
LIVER FUNCTION TEST (LFT) (S)			
Alkaline Phosphatase IFCC PNPP Buffer	138.7	U/L	44 - 147 U/L
Total Protein BIURET	6.8	gm/dl	6.0 - 8.3
Albumin BCG	3.98	g/dl	3.5 - 5.5 g/dl
Globulin	2.82	gm/dl	2.0 - 3.5 gm/dl
AG RATIO	1.16		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 CHOD - PAP	219.7	mg/dl	<200.0 mg/dl
Triglycerides GPO - PAP	159.7	mg/dl	< 150 mg/dl
HDL Cholesterol Homogeneous Enzymatic Colorimetric test	44.31	mg/dl	Adult males >45 mg/dl
LDL Cholesterol	143.45	mg/dl	<100 mg/dl
VLDL Cholesterol	31.94	mg/dl	<30.0 mg/dl
CHO/HDL Ratio	4.96	mg/dl	Low risk 3.3-4.4
Non HDL Cholesterol Calculated	175.39	mg/dl	<130 mg/dl

Sample Type : SERUM



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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 30 ml
pH 6.0
Mucus ABSENT
Absent, Present
Appearance CLEAR
Slightly turbid, Turbid, Clear

Chemical Examination (Strip)

Specific Gravity 1.015
Albumin TRACE
Absent, Present(+), Present(2+), Present(3+)
Sugar 4(+)
Absent, Present(+), Present(2+), Present(3+)
Bilirubin NEGATIVE
Absent, Present

Microscopic Examination (Microscopy)

Pus Cells 6-8 /HPF
Epithelial Cells 2-4 /HPF
RBC NIL /HPF
Casts ABSENT
Crystals ABSENT
Bacteria ABSENT

Others
Sample Type : Urine

Laboratory

Glucose, Fasting 275.1 mg/dl 70 - 110 mg/dl
Sample Type : SERUM



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- OR
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Note:
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ENDOCRINE

Thyroid Hormones (T3 .T4 & TSH)

T3	0.72	ng/ml	0.60 - 1.81 ng/ml
T4	8.13	ng/dl	5.01 - 12.45 ng/dl
TSH (Thyroid stimulating hormones)	5.11	uIU/ml	0.34 - 5.50 uIU/ml

Sample Type : SERUM



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Remarks :

Note1. TSH levels are subject to circadian variation, reaching peak levels between 2-4.a.m and at a minium 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
- * Hperthyroidism
- * Hypothalamic- Pituitary hypothyroidism
- * Inappropriate-TSH secretion
- * Nonthyroidal illness
- * Autoimmune thyroid disease
- * Pregnancy associated thyroid disorders
- * Thyroid dysfunction in infancy and early childhood

IMMUNOLOGY

Total PSA	0.89	ng/ml	0.00 - 4.0 ng/ml
Sample Type :	SERUM		

Summary & Interpretation:

Elevated concentrations of PSA in serum are generally indicative of a patho-logic-condition of the prostate (prostatitis, begin hyperplasia or carcinoma). PSA determinations are employed are the monitoring of progress and efficiency of therapy in patients with prostate carcinoma or receiving hormonal therapy . An inflammation or trauma of the prostate(e.g. In case of urinary retention or following rectal examination, cystoscopy, coloscopy, transurethral biopsy, lasertreatment or ergometry) can lead to PSA elevations of varying duration and magnitu

--End of Report--