



Name : Mrs. NEENA W/o UHID : 120386 S No : PID : 33295
Age/Gender : 45 Year/Female A.S : NP Sample Date : 31-Jul-2024 11:16 AM
Ref. By Dr. : MEDIWHEEL Report Date : 31-Jul-2024 05:37 PM
Address : HISAR Status : Pending Sample Type : Inside *33295*

Test Name Value Unit Reference Range

HEMATOLOGY

CBC (Complete Blood Count)

Haemoglobin (Hb)	13.0	g/dl	12.0 - 15.0 g/dl
Total RBC Count	4.36	m/cumm	4.20 - 5.40
Haematocrit	39.0	%	35.0 - 50.0 %
Mean Cell Volume	89.4	fL	80.0 - 100 fL
Mean Cell Haemoglobin	29.8	pg	27.0 - 34.0 pg
Mean Cell Haemoglobin Conc	33.4	%	32.0 - 36.0
Red Cell Distribution Width (RDW)-CV	12.6	%	11.0 - 16.0 %
Red Cell Distribution Width (RDW)-SD	45.7	fL	35.0 - 56.0 fL
Total Leucocyte Count	6600	cells/cum m	4000 - 11000
Differential Leucocyte Count	.		
Neutrophils	50	%	32 - 72 %
Lymphocytes	45	%	20 - 50 %
Monocytes	03	%	2 - 11 %
Eosinophils	02	%	1 - 3 %
Basophils	0	%	0 - 2 %
Platelet Count	2,97,000	cells/cum m	150,000 - 450,000
Platelet Distribution Width	15.3	fL	15.0 - 18.0 fL
Mean Platelet Volume	9.7	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

O⁺ 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 5.6 % 4.27 - 6.00 %



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Test Name	Value	Unit	Reference Range
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HBA1C

turbidimetric immunoassay

Average Blood Glucose

114.02

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	80	mmHr	0 - 20 mmHr
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Sample Type : Whole Blood



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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	4.01	mg/dL	2.5 - 6.0
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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.

Total Protein

Total Protein	6.9	gm/dl	6.0 - 8.3
BIURET			
Albumin	4.19	g/dl	2.9 - 4.5
BCG			
Globulin	2.71	gm/dl	2.0 - 3.5
Albumin-Globulin Ratio	1.48		1.2 - 2.5



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Test Name	Value	Unit	Reference Range
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Sample Type : SERUM

UREA. SERUM

UREA	25.89	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	0.9	mg/dL	0.5 - 1.4 mg/dL
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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.

KIDNEY FUNCTION TEST (KFT Special)

UREA	25.89	mg/dL	14 - 45 mg/dL
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KINETIC METHOD WITH UREASE AND GLDH

CREATININE SERUM

Jaffe Kinetic

Uric acid

Uricase - POD

BUN SERUM

KINETIC METHOD WITH UREASE & GLDH

SODIUM-SERUM

ISE(DIRECT)

POTASSIUM SERUM

ISE(DIRECT)

Chloride

Ion Selective Electrode (indirect)

Urea / Creatinine Ratio

BUN / Creatinine Ratio

Sample Type : SERUM



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Test Name	Value	Unit	Reference Range
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CLINICAL COMMENTS :

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: 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.

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.

SODIUM: Increases due to water loss (severe diarrhea profuse sweating, polyuria or vomiting), hypergluco- or mineralo-corticoidism, and inadequate water intake. Decreases due to intake of free water or

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	19.70	IU/L	10 - 40 IU/L
IFCC with Pyridoxal Phosphate			
SGPT	27.10	IU/L	07 - 56 IU/L
IFCC with Pyridoxal Phosphate			
Alkaline Phosphatase	62.08	U/L	44 - 147 U/L
IFCC PNPP Buffer			
Total Protein	6.9	gm/dl	6.0 - 8.3
BIURET			
Albumin	4.19	g/dl	3.5 - 5.5 g/dl
BCG			
Globulin	2.71	gm/dl	2.0 - 3.5 gm/dl
AG RATIO	1.48		1.2 - 2.5

Sample Type : SERUM



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Test Name	Value	Unit	Reference Range
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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	158.40	mg/dl	<200.0 mg/dl
CHOD - PAP			
Triglycerides	250.4	mg/dl	< 150 mg/dl
GPO - PAP			
HDL Cholesterol	41.8	mg/dl	Adult females >55 mg/dl
Homoogeneous Enzymatic Colorimetric test			
LDL Cholesterol	66.52	mg/dl	<100 mg/dl
VLDL Cholesterol	50.08	mg/dl	<30.0 mg/dl
CHO/HDL Ratio	3.79	mg/dl	Low risk 3.3-4.4
Non HDL Cholesterol	116.6	mg/dl	<130 mg/dl

Calculated

Sample Type : SERUM

Interpretation

Note

1. 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.

2. 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.

3. Apolipoprotein B is an optional, secondary lipid target for treatment once LDL & Non HDL goals have been achieved.

4. 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



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Test Name	Value	Unit	Reference Range
Colour Pale-yellow, Yellowish, Colorless, YELLOW	PALE YELLOW		
Quantity	40	ml	
pH	7.0		
Mucus Absent, Present	ABSENT		
Appearance Slightly turbid, Turbid, Clear	CLEAR		
Chemical Examination (Strip)	.		
Specific Gravity	1.025		
Albumin Absent, Present(+), Present(2+), Present(3+)	NEGATIVE		
Sugar Absent, Present(+), Present(2+), Present(3+)	NEGATIVE		
Bilirubin Absent, Present	NEGATIVE		
Microscopic Examination (Microscopy)	.		
Pus Cells	2-4	/HPF	
Epithelial Cells	1-2	/HPF	
RBC	NIL	/HPF	
Casts	ABSENT		
Crystals	ABSENT		
Bacteria	ABSENT		
Others			
Sample Type : Urine			

Laboratory

Blood Sugar (PP) Blood Sugar PP Sample Type : Others	96.60	mg/dl	70.00 - 140.00 mg/dl
Glucose, Fasting Sample Type : SERUM	77.1	mg/dl	70 - 110 mg/dl



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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 ≥ 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 .

URINE SUGAR FASTING	79.8		
Sample Type : Urine			
URINE SUGAR PP	79.8		70 - 110
Sample Type : Urine			

ENDOCRINE

Thyroid Hormones (T3 .T4 & TSH)

T3	1.13	ng/ml	0.60 - 1.81 ng/ml
T4	10.24	ng/dl	5.01 - 12.45 ng/dl
TSH Ultrasensitive	2.26	uIU/ml	0.3 - 4.5 uIU/ml
Sample Type : SERUM			



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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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--

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MBBS, MD
Consultant Pathologist

Print Date : 7/31/2024 10:15

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PATIENT NAME: NEENA
REF BY: TPA

AGE/SEX: 45 YRS/F
DATE: JULY 31, 2024

USG WHOLE ABDOMEN

Liver: normal in size and **shows mild fatty changes**. No focal area of altered echogenicity is seen. IHBR not dilated. Portal vein is normal.

Gall bladder: is not visualized (H/o Cholecystectomy). CBD measures 6.5 mm.

Pancreas: head and body shows normal size and parenchymal attenuation.

Spleen: normal in size and normal echotexture.

Right Kidney: is normal in position, size and morphology. No evidence of any calculus detected. Pelvi calyceal system is normal. CMD is maintained.

Left Kidney: is normal in position, size and morphology. No evidence of any calculus detected. Pelvi calyceal system is normal. CMD is maintained.

Urinary Bladder: appears normal.

Uterus: is not visualized (H/o hysterectomy).

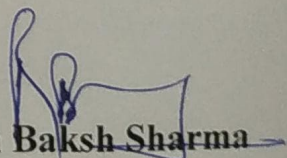
No adnexal mass lesion seen.

No obvious abnormal bowel dilatation or wall thickening is seen in present scan.

No free fluid seen.

IMPRESSION: - Mild fatty changes in liver.

Clinical correlation and further evaluation is suggested.


Dr. Ram Baksh Sharma
Radiologist

Dr. Rambaksh Sharma
Consultant Radiologist

Dr. Anshul Jain
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Dr. Rajesh Reddu
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Dr. Sonam Aneja
Consultant Pathologist

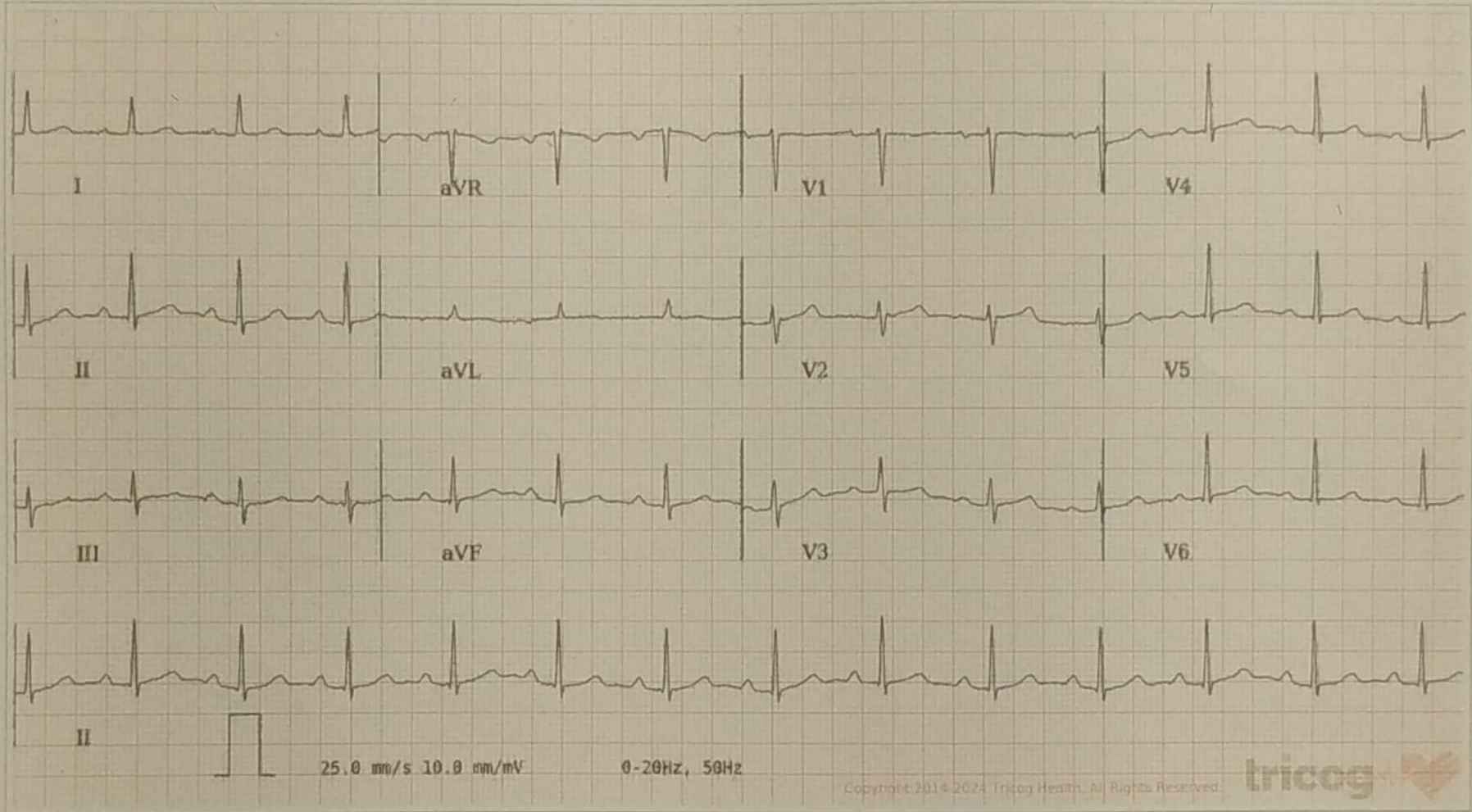


GEETANJALI HOSPITAL HISAR HARYANA I

Age / Gender: 45/Female
Patient ID: 0000312460

Date and Time: 31st Jul 24 12:45 PM

Mrs Meena



AR: 84bpm VR: 83bpm QRSD: 72ms QT: 346ms QTcB: 409ms PR: 210ms P-R-T: 56° 37° 53°

Sinus Rhythm, First Degree AV Block. Please correlate clinically.

REPORTED BY

Nisar Ahmad K
Dr. Nisar Ahmad K
KMC 122453