



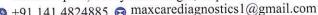


Sylym

Dr. U. G. GUPTA MBBS, MD (Physician) RMC No. 291











Date of Examination: 2412
Name: MAHESH CHAND YADAY Age: 34 YRSDOB: 06/06/1988Sex: MALE
Referred By: BANKOFBARODA
Photo ID: TDCARD ID#: 92882
Ht: 183 (cm) Wt: 3 (Kg)
Chest (Expiration): <u>98</u> (cm) Abdomen Circumference: <u>99</u> (cm)
Blood Pressure: 124 03 mm Hg PR: 78 min RR: 10 min Temp: Afelonic
BMI 22
Eye Examination: RIET G/G, N/G, NCB LIEJ G/G, N/G, NCB
Other: N/A
On examination he/she appears physically and mentally fit: Yes/No
Signature Of Examine: MAHESH.CHAND YADAY
Signature Medical Examiner: Name Medical Examiner - UoC CIUPTA
MBBS, MD (Physician) RMC No. 291



P3 HEALTH SOLUTIONS LLP

(ASSOCIATES OF MAXCARE DIAGNOSTICS)

 B-14, Vidhyadhar Enclave - II, Near Axis Bank Central Spine, Vidhyadhar Nagar, Jaipur - 302023

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NAME :- Mr. MAHESH CHAND YADAV

Age:- 34 Yrs 6 Mon 20 Days

Sex :- Male



Patient ID :-12222722 Date :- 24/12/2022

Ref. By Doctor:-BANK OF BARODA

Lab/Hosp :-

Company :- Mr.MEDIWHEEL

Final Authentication: 24/12/2022 18:49:00

09:36:52

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP BELOW 40	MALE .		
	IVIALL		
HAEMOGARAM	140		12.0 17.0
HAEMOGLOBIN (Hb)	14.9	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	3.40 └	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	52.0	%	40.0 - 80.0
LYMPHOCYTE	40.0	%	20.0 - 40.0
EOSINOPHIL	3.0	%	1.0 - 6.0
MONOCYTE	5.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	4.87	x10^6/uL	4.50 - 5.50
HEMATOCRIT (HCT)	47.50	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	98.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	30.6	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	31.3 L	g/dL	31.5 - 34.5
PLATELET COUNT	169	x10^3/uL	150 - 410
RDW-CV	14.2 H	%	11.6 - 14.0
MENTZER INDEX	20.12 H		0.00 - 0.00

A complete blood picture (CBP) is a kind of blood test that is done to assess a person's overall health and diagnose a wide range of health disorders like leukemia, anemia and other infections.

A complete blood count (CBC) is a complete blood test that diagnose many components and features of a persons blood which includes: -

(CBC): Methodology: TLC,TRBC,PCV,PLT Impedance method, HB Calorimetric method, and MCH,MCV,MCHC,MENTZER INDEX are calculated. InstrumentName: MINDRAY BC-3000 Plus 3 part automatic analyzer,

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MD (Pathology) RMC No. 17226

Janu

^{*}Red Blood Cells (RBC), which carry oxygen -

^{*}White Blood Cells (WBC), which help in fighting against infections -

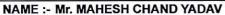
^{*}Hemoglobin, which is the oxygen carrying protein in the red blood cells -

^{*}Hematocrit (HCT), the proportion of RBC to the fluid component, or plasma present in blood -

^{*}Platelets, which aid in blood clotting



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HAEMATOLOGY

Erythrocyte Sedimentation Rate (ESR)

Methord:- Westergreen

07

mm in 1st hr

00 - 15

The erythrocyte sedimentation rate (ESR or sed rate) is a relatively simple, inexpensive, non-specific test that has been used for many years to help detect inflammation associated with conditions such as infections, cancers, and autoimmune diseases.ESR is said to be a non-specific test because an elevated result often indicates the presence of inflammation but does not tell the health practitioner exactly where the inflammation is in the body or what is causing it. An ESR can be affected by other conditions besides inflammation. For this reason, the ESR is typically used in conjunction with other tests, such as C-reactive protein.ESR is used to help diagnose certain specific inflammatory diseases, including temporal arteritis, systemic vasculitis and polymyalgia rheumatica. (For more on these, read the article on Vasculitis.) A significantly elevated ESR is one of the main test results used to support the diagnosis. This test may also be used to monitor disease activity and response to therapy in both of the above diseases as well as



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NAME :- Mr. MAHESH CHAND YADAV 34 Yrs 6 Mon 20 Days

Sex :-Male

Test Name

Age :-

Company :-

BIOCHEMISTRY

Value Unit **Biological Ref Interval**

FASTING BLOOD SUGAR (Plasma) Methord:- GOD POD

81.1

mg/dl

70.0 - 115.0

Impaired glucose tolerance (IGT) Diabetes Mellitus (DM)

111 - 125 mg/dL

> 126 mg/dL

Instrument Name: HORIBA CA60 Interpretation: Elevated glucose levels (hyperglycemia) may occur with diabetes, pancreatic neoplasm,

hyperthyroidism and adrenal cortical hyper-function as well as other disorders, Decreased glucose levels(hypoglycemia) may result

from excessive insulin

therapy or various liver diseases.



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DR.TANU RUNGTA MD (Pathology)

RMC No. 17226

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Final Authentication: 25/12/2022 11:52:24

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval

FULL BODY HEALTH CHECKUP BELOW 40 MALE

BLOOD SUGAR PP (Plasma) Methord:- GOD PAP 97.4

mg/dl

70.0 - 140.0

Instrument Name: HORIBA Interpretation: Elevated glucose levels (hyperglycemia) may occur with diabetes, pancreatic neoplasm, hyperthyroidism and adrenal cortical hyper-function as well as other disorders. Decreased glucose levels (hypoglycemia) may result from excessive insulin therapy or various liver diseases.



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Page No: 1 of 1

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34 Yrs 6 Mon 20 Days Age :-

Male Sex :-



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HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval	
GLYCOSYLATED HEMOGLOBIN (Hb.A. Methord:- CAPILLARY with EDTA	5.3	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0	
MEAN PLASMA GLUCOSE Methord:- Calculated Parameter	102	mg/dL	68 - 125	

INTERPRETATION

AS PER AMERICAN DIABETES ASSOCIATION (ADA) Reference Group HbA1c in % Non diabetic adults >=18 years < 5.7 At risk (Prediabetes) 5.7 - 6.4 Diagnosing Diabetes >= 6.5

CLINICAL NOTES

In vitro quantitative determination of HbA1c in whole blood is utilized in long term monitoring of glycemia. The HbA1c level correlates with the mean glucose concentration prevailing in the course of the patient's recent history (approx - 6-8 weeks) and therefore provides much more reliable information for glycemia monitoring than do determinations of blood glucose or urinary glucose. It is recommended that the determination of HbA1c be performed at intervals of 4-6 weeks during Diabetes Mellitus therapy. Results of HbA1c should be assessed in conjunction with the patient's medical history, clinical examinations and other findings. Some of the factors that influence HbA1c and its measurement [Adapted from Gallagher et al]

1. Erythropoiesis

- Increased HbA1c: iron, vitamin B12 deficiency, decreased erythropolesis.
- Decreased HbA1c: administration of erythropoietin, iron, vitamin B12, reticulocytosis, chronic liver disease
- 2. Altered Haemoglobin-Genetic or chemical alterations in hemoglobin: hemoglobinopathies, HbF, methemoglobin, may increase or decrease HbA1c.

- Increased HbA1c: alcoholism, chronic renal failure, decreased intraerythrocytic pH
- Decreased HbA1c: certain hemoglobinopathies, increased intra-erythrocyte pH

4. Erythrocyte destruction

- Increased HbA1c; increased erythrocyte life span; Splenectomy
- Decreased A1c: decreased RBC life span; hemoglobinopathies, splenomegaly, rheumatoid arthritis or drugs such as antiretrovirals, ribavirin & dapsone.

- Increased HbA1c: hyperbilirubinemia, carbamylated hemoglobin, alcoholism, large doses of aspirin, chronic opiate use chronic renal failure
- Decreased HbA1c: hypertriglyceridemia, reticulocytosis, chronic liver disease, aspirin, vitamin C and E, splenomegaly, rheumatoid arthritis or drugs

1. Shortened RBC life span - HbA1c test will not be accurate when a person has a condition that affects the average lifespan of red blood cells (RBCs), such as hemolytic anemia or blood loss. When the lifespan of RBCs in circulation is shortened, the A1c result is falsely low and is an unreliable measurement of a person's average glucose over time.

2. Abnormal forms of hemoglobin – The presence of some hemoglobin variants, such as hemoglobin S in sickle cell anemia, may affect certain methods for measuring A1c. In these cases, fructosamine can be used to monitor glucose control.

Advised:

1.To follow patient for glycemic control test like fructosamine or glycated albumin may be performed instead.

2. Hemoglobin HPLC screen to analyze abnormal hemoglobin variant.

estimated Average Glucose (eAG) : based on value calculated according to National Glycohemoglobin Standardization Program (NGSP) criteria

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HAEMATOLOGY

BLOOD GROUP ABO Methord:- Haemagglutination reaction "B" POSITIVE



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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE TOTAL CHOLESTEROL Methord:- CHOD-PAP methodology	118.00	mg/dl	Desirable <200 Borderline 200-239 High> 240
InstrumentName: MISPA PLUS Interpretation: disorders.	Cholesterol measurement	s are used in the diagnosis a	and treatments of lipid lipoprotein metabolism
TRIGLYCERIDES Methord:- GPO-TOPS methodology	51.20	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500

InstrumentName: MISPA PLUS Interpretation: Triglyceride measurements are used in the diagnosis and treatment of diseases involving lipid metabolism and various endocrine disorders e.g. diabetes mellitus, nephrosis and liver obstruction.

DIRECT HDL CHOLESTEROL Methord:- Selective inhibition Method

mg/dl

Male 35-80 Female 42-88

Instrument Name: MISPA PLUS Interpretation: An inverse relationship between HDL-cholesterol (HDL-C) levels in serum and the incidence/prevalence of coronary heart disease (CHD) has been demonstrated in a number of epidemiological studies. Accurate measurement of HDL-C is of vital importance when assessing patient risk from CHD. Direct measurement gives improved accuracy and reproducibility when compared to precipitation methods 44 07

LDL CHOLESTEROL Methord:- Calculated Method

mg/dl Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190 mg/dl 0.00 - 80.00

Methord:- Calculated T.CHOLESTEROL/HDL CHOLESTEROL RATIO 1.80 Methord: - Calculated

LDL / HDL CHOLESTEROL RATIO Methord:- Calculated

0.67

10.24

0.00 - 4.90

0.00 - 3.50

TOTAL LIPID Methord:- CALCULATED

VLDL CHOLESTEROL

336.50 L

mg/dl

400.00 - 1000.00

1. Measurements in the same patient can show physiological & analytical variations. Three serialsamples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL& LDL Cholesterol.

- 2. As per NCEP guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended
- 3. Low HDL levels are associated with Coronary Heart Disease due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated fromperipheral tissues.

Comments: 1- ATP III suggested the addition of Non HDL Cholesterol (Total Cholesterol - HDL Cholesterol) as an indicator of all VIKARANTJI

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Sex :-



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BIOCHEMISTRY

LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Methord:- DMSO/Diazo	0.57	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Methord:- DMSO/Diazo	0.23	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Methord:- Calculated	0.34	mg/dl	0.30-0.70
SGOT Methord:- IFCC	40.0 H	U/L	Men- Up to - 37.0 Female - Up to - 31.0
SGPT Methord:- IFCC	38.4	U/L	Men- Up to - 40.0 Female- Up to - 31.0
SERUM ALKALINE PHOSPHATASE Methord:- DGKC - SCE	57.60	U/L	53.00 - 141.00
SERUM GAMMA GT Methord: - Szasz methodology Instrument Name Randox Rx Imola Interpretation: Elevations in GGT levels are seen earlier and more pronounced than	19.20 a those with other liver enz	U/L ymes in cases of obstructive jaundice and	10.00 - 45.00
metastatic neoplasms. It may reach 5 to 30 times normal levels in intra-or post- hepatic biliary obstruction. Only moderate elevations in the enzyme level (2 to 5 ti	mes normal)are observed v	with infectious hepatitis.	
SERUM TOTAL PROTEIN Methord:- Direct Biuret Reagent	6.83	g/dl	5.10 - 8.00
SERUM ALBUMIN Methord:- Bromocresol Green	4.87	g/dl	3.50 - 5.50
SERUM GLOBULIN Methord:- CALCULATION	1.96 └	gm/dl	2.20 - 3.50
The state of the s	COMMISSION	A STATE OF THE PARTY OF THE PAR	

Interpretation: Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

Note:- These are group of tests that can be used to detect the presence of liver disease, distinguish among different types of liver disorders, gauge the extent of known liver damage, and monitor the response to treatment. Most liver diseases cause only mild symptoms initially, but these diseases must be detected early. Some tests are associated with functionality (e.g., albumin), some with cellular integrity (e.g., transaminase), and some with conditions linked to the biliary tract (gamma-glutamyl transferase and alkaline phosphatase). Conditions with elevated levels of ALT and AST include hepatitis A,B,C, paracetamol toxicity etc. Several biochemical tests are useful in the evaluation and management of patients with hepatic dysfunction. Some or all of these measurements are also carried out (usually about twice a year for routine eases) on those individuals taking certain medications, such as anticonvulsants, to ensure that the medications are not adversely impacting the person's liver.

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A/G RATIO

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DR.TANU RUNGTA

1.30 - 2.50

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BIOCHEMISTRY

RFT / KFT WITH ELECTROLYTES

SERUM UREA Methord:- Urease/GLDH 19.60

mg/dl

10.00 - 50.00

InstrumentName: HORIBA CA 60 Interpretation: Urea measurements are used in the diagnosis and treatment of certain renal and metabolic

diseases.

SERUM CREATININE Methord:- Jaffe's Method

1.26

mg/dl

Males: 0.6-1.50 mg/dl

Females: 0.6 -1.40 mg/dl

Interpretation:

Creatinine is measured primarily to assess kidney function and has certain advantages over the measurement of urea. The plasma level of creatinine is relatively independent of protein ingestion, water intake, rate of urine production and exercise. Depressed levels of plasma creatinine are rare and not

clinically significant. SERUM URIC ACID

4.42

mg/dl

2.40 - 7.00

InstrumentName: HORIBA YUMIZEN CA60 Daytona plus Interpretation: Elevated Urate: High purine diet, Alcohol. Renal insufficiency, Drugs, Polycythaemia vera, Malignancies, Hypothyroidism, Rare enzyme defects, Downs syndrome, Metabolic syndrome, Pregnancy, Gout.

SODIUM Methord:- ISE

Interpretation: Decreased sodium - Hyponatraemia Causes include: fluid or electrolyte loss, Drugs, Oedematous states, Legionnaire's disease and other chest infections, pseudonatremia, Hyperlipidaemias and paraproteinaemias, endocrine diseases, SIADH.

POTASSIUM

Methord: - ISE

4.53

mmol/L

3.50 - 5.50

Interpretation: A. Elevated potassium (hyperkalaemia). Artefactual, Physiological vation, Drugs, Pathological states, Renal failure Adrenocortical insufficiency, metabolic acidoses, very high platelet or white cell counts B. Decreased potassium (hypokalaemia)Drugs, Liquoric, Diarrhoea and vomiting, Metabolic alkalosis, Corticosteroid excess, Oedematous state, Anorexia nervosa/bulimia

CHLORIDE

101.4

mmol/L

94.0 - 110.0

Interpretation: Used for Electrolyte monitoring.

SERUM CALCIUM

Methord:- Arsenazo III Method

9.98

mg/dL

8.80 - 10.20

InstrumentName:MISPA PLUS Interpretation: Serum calcium levels are believed to be controlled by parathyroid hormone and vitamin D. Increases in serum PTH or vitamin D are usually associated with hypercalcemia .Hypocalcemia may be observed in hypoparathyroidism, nephrosis and pancreatitis.

SERUM TOTAL PROTEIN VNGARIA NiFef Biuret Reagent

6.83

g/dl

5.10 - 8.00 Janu

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DR.TANU RUNGTA

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BIOCHEMISTRY

SERUM ALBUMIN Methord:- Bromocresol Green 4.87

g/dl

3.50 - 5.50

SERUM GLOBULIN

1.96 L

gm/dl

2.20 - 3.50

A/G RATIO

2.48

1.30 - 2.50

Interpretation: Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

INTERPRETATION

Kidney function tests are group of tests that can be used to evaluate how well the kidneys are functioning. Creatinine is a waste product that comes from protein in the diet and also comes from the normal wear and tear of muscles of the body. In blood, it is a marker of GFR .in urine, it can remove the need for 24-hour collections for many analytes or be used as a quality assurance tool to assess the accuracy of a 24-hour collection Higher levels may be a sign that the kidneys are not working properly. As kidney disease progresses, the level of creatinine and urea in the bloodincreases. Certain drugs are nephrotoxic hence KFT is done before and after initiation of treatment with these drugs.

Low serum creatinine values are rare; they almost always reflect low muscle mass.

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Janu DR.TANU RUNGTA MD (Pathology)

RMC No. 17226



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TOTAL THYROID PROFILE

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
THYROID-TRIIODOTHYRONINE T3 Methord:- ECLIA	1.12	ng/mL	0.70 - 2.04

NOTE-TSH levels are subject to circardian variation, reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration. Dose and time of drug intake also influence the test result. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions, simoultaneous measurement of TSH with free T4 is useful in evaluating differential diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay 1. Primary hyperthyroidism is accompanied by † serum T3 & T4 values along with * TSH level. 2. Low TSH, high FT4 and TSH receptor antibody(TRAb) +ve seen in patients with Graves disease 3.Low TSH high FT4 and TSH receptor antibody (TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter 4.HighTSH.Low FT4 and Thyroid microsomal +ve seen in patients with Graves disease 3.Low TSH,high F14 and TSH receptor antibody (TRAb) -ve seen in patients with Toxic adenomal Toxic Multinodular golder 4.High TSH,Low F14 and TSH receptor antibody increased seen in patients with Hashimotos thyroiditis 5.HighTSH,Low F14 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency 6.Low TSH,Low F14 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism 7.Primary hypothyroidism is accompanied by ‡ serum T3 and T4 values & 'serum TSH levels 8.Normal T4 levels accompanied by *T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis9.Normal or T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .11.Normal T3 & *T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild / Subclinical Hypothyroidism .15.Normal T3 & T4 along with *TSH indicate mild *TSH in

DURING PREGNANCY - REFERENCE RANGE for TSH IN uIU/mL (As per American Thyroid Association) 1st Trimester: 0.10-2.50 uIU/mL 2nd Trimester: 0.20-3.00 uIU/mL 3rd Trimester: 0.30-3.00 ulU/mL The production, circulation, and disintegration of thyroid hormones are altered throughout the stages of pregnancy

REMARK-Assay results should be interpreted in context to the clinical condition and associated results of other investigations. Previous treatment with corticosteroid therapy may result in lower TSH levels while thyroid hormone levels are normal. Results are invalidated if the client has undergone a radionuclide scan within 7-14 days before the test. Abnormal thyroid test findings often found in critically ill patients should be repeated after the critical nature of the condition is resolved. TSH is an important marker for the diagnosis of thyroid dysfunction. Recent studies have shown that the TSH distribution progressively shifts to a higher phyrroid dysfunction of the condition is resolved. TSH is an important marker for the diagnosis of thyroid dysfunction. Recent studies have shown that the TSH distribution progressively shifts to a higher phyrroid disease in the elderly. The condition is resolved. TSH is an important marker for the diagnosis of thyroid dysfunction. Recent studies have shown that the TSH distribution progressively shifts to a higher phyrroid disease in the elderly. The condition is resolved. The condition is res

NOTE-TSH levels are subject to circardian variation, reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration. Does and time of drug intake also influence the test result. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions, simoultaneous measurer of TSH with free T4 is useful in evaluating differential diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay 1. Primary hyperthyroidism is accompanied by tserum T3 & T4 values along with * TSH level. 2. Low TSH, high FT4 and TSH receptor antibody (TRAb) IN LCR RC IATION-Ultra Sensitive 4th generation assay 1. Phimary hypothyroidism is accompanied by fserum 13 & 14 values along with "TSH levels along with "TSH levels along with "14 and TSH receptor antibody (TRAb) -ve seen in patients with Toxic Multinodular goiter 4. HighTSH, Low FT4 and TNyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5. HighTSH, Low FT4 and TNyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency 6. Low TSH, Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertlary hypothyroidism
7. Primary hypothyroidism is accompanied by 1 serum T3 and T4 values & 'serum TSH levels 8. Normal T4 levels accompanied by "T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis9 Normal or "T3 & "T4 along with" TSH indicate mild / Subclinical Hypoth

DURING PREGNANCY - REFERENCE RANGE for TSH IN uIU/mL (As per American Thyroid Association) 1st Trimester: 0.10-2.50 uIU/mL 2nd Trimester: 0.20-3.00 uIU/mL 3rd Trimester: 0.30-3.00 uIU/mL. The production, circulation, and disintegration of thyroid hormones are altered throughout the stages of pregnancy.

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TSH Methord:- ECLIA 5.266

μIU/mL

0.350 - 5.500

NOTE-TSH levels are subject to circardian variation, reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration.Dose and time of drug intake also influence the test result.

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NTERPRETATION-Ultra Sensitive 4th generation assay

Technologist

Page No: 14 of 15

Janu DR.TANU RUNGTA MD (Pathology) RMC No. 17226



(ASSOCIATES OF MAXCARE DIAGNOSTICS)

 B-14, Vidhyadhar Enclave - II, Near Axis Bank Central Spine, Vidhyadhar Nagar, Jaipur - 302023

© +91 141 4824885 maxcarediagnostics1@gmail.com

NAME :- Mr. MAHESH CHAND YADAV 34 Yrs 6 Mon 20 Days

Sex :-Male

Age :-

Patient ID: -12222722 Date :- 24/12/2022

Ref. By Doctor:-BANK OF BARODA

Lab/Hosp:-

Company :-Mr.MEDIWHEEL

Final Authentication: 24/12/2022 18:49:00

09:36:52

CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
Urine Routine PHYSICAL EXAMINATION			
COLOUR	PALE YEL	LOW	PALE YELLOW
APPEARANCE	Clear		Clear
CHEMICAL EXAMINATION			
REACTION(PH)	5.0		5.0 - 7.5
SPECIFIC GRAVITY	1.030		1.010 - 1.030
PROTEIN	NIL	A.	NIL
SUGAR	NIL	THE STATE OF THE S	NIL
BILIRUBIN	NEGATIVI	Ε	NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVI	Ε , , , , ,	NEGATIVE
NITRITE	NEGATIVE		NEGATIVE
MICROSCOPY EXAMINATION	een Y		
RBC/HPF	NIL	/HPF	NIL
WBC/HPF	2-3	/HPF	2-3
EPITHELIAL CELLS	2-3	/HPF	2-3
CRYSTALS/HPF	ABSENT		ABSENT
CAST/HPF	ABSENT		ABSENT
AMORPHOUS SEDIMENT	ABSENT		ABSENT
BACTERIAL FLORA	ABSENT		ABSENT
YEAST CELL	ABSENT		ABSENT
OTHER	ABSENT		

VIKARANTJI

Technologist

Page No: 12 of 15

Janu DR.TANU RUNGTA

MD (Pathology) RMC No. 17226



O B-14, Vidhyadhar Enclave - II, Near Axis Bank Central Spine, Vidhyadhar Nagar, Jaipur - 302023 9 +91 141 4824885 maxcarediagnostics1@gmail.com



NAME:	MR. MAHESH CHAND YADAV	AGE	34 YRS/M
REF.BY	BANK OF BARODA	DATE	24/12/2022

CHEST X RAY (PA VIEW)

Bilateral lung fields appear clear.

Bilateral costo-phrenic angles appear clear.

Cardiothoracic ratio is normal.

Thoracic soft tissue and skeletal system appear unremarkable.

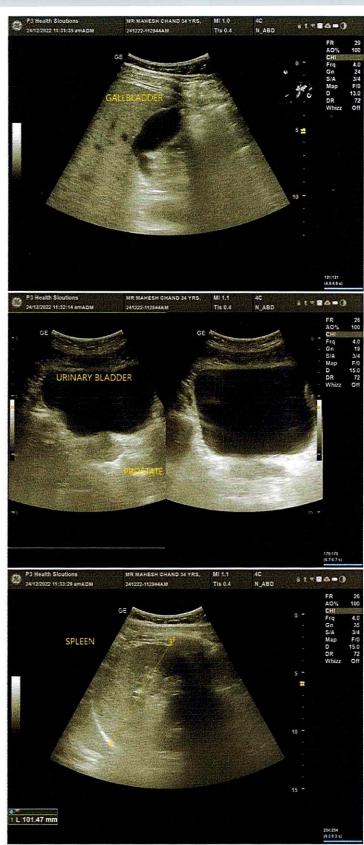
Soft tissue shadows appear normal.

IMPRESSION: No significant abnormality is detected.

Shallni

DR.SHALINI GOEL M.B.B.S, D.N.B (Radiodiagnosis) RMC No.: 21954







🕲 +91 141 4824885 😂 maxcarediagnostics1@gmail.com



MR. MAHESH CHAND YADAV	34 Y/Male
Registration Date: 24/12/2022	Ref. by: MANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (13.5 cm). **Echotexture is increased obscuring periportal echogenicity.** No focal space occupying lesion is seen within liver parenchyma. Intra hepatic biliary channels are not dilated. Portal vein diameter is normal.

Gall bladder is well distended. No calculus or mass lesion is seen in gall bladder. Common bile duct is not dilated.

Pancreas is of normal size and contour. Echo-pattern is normal. No focal lesion is seen within pancreas.

Spleen is of normal size and shape (10.1 cm). Echotexture is normal. No focal lesion is seen.

Kidneys are normally sited and are of normal size and shape. Cortico-medullary echoes are normal. Collecting system does not show any calculus or dilatation.

Right kidney is measuring approx. 9.9 x 4.0 cm.

Left kidney is measuring approx. 11.7 x 4.9 cm.

Urinary bladder does not show any calculus or mass lesion.

Prostate is normal in size with normal echotexture and outline.

No enlarged nodes are visualized. No retro-peritoneal lesion is identified. No significant free fluid is seen in pelvis.

IMPRESSION:

- Grade-2 fatty liver.
- · Rest no significant abnormality is detected.

Shallni

DR.SHALINI GOEL M.B.B.S, D.N.B (Radiodiagnosis) RMC no.: 21954

MBBS, DNB (Radiologist)
RMC No. 21954
P-3 Health Solutions LLP

lef.: BANK OF BARODA 12229451322741/Mr Mahesh Chand Yadav 34Yrs/Male Comments: P-QRS-T axis: 60. 23. 81. (Deg) Vent Rate: 50 bpm; PR Interval: 108 ms; FINDINGS: Normal Sinus Rhythm avR Test Date: 24-Dec-2022(12:25:42) Notch: 50Hz 0.05Hz - 100Hz QRS Duration: Kgs/ Cms avL Y 128 ms; QT/QTc Int: 411/377 10mm/mV 25mm/Sec mmHg HR: 50 bpm Basoleswine 8 5 హ PR Interval: 108 ms QRS Duration: 128 ms QT/QTc: 411/377ms P-QRS-T Axis: 60 - 23 - 81 (Deg) Dr. Naresh Kunga-Mohanka RMCAO:: 35703 ABBS, DIP. CARDIO (ESCORTS) D.E.M. (RCGP-UK) which beginsty and a

3-14, Vidhyanagar Nagar, Enclave, Phase-2, Jaipur

BP:

3 HEALIH SULULIUNS LLF

Protocol : BRUCE

ate: 24-Dec-2022 12:27:23 PM ef.By : BANK OF BARODA 322304/MR MAHESH CHAND YADAV 34 Yrs/Male 0 Kg/0 Cms i-14, Vidhyadhar Nagar Enclave, Phase -2, Jaipur

tage 2 'eakEx dvice/Comments: indings ecovery ecovery upine tage ecovery ecovery tage 1 edication: xStart tanding bjective: Max BP : 160/90(mmHg) Max HR Attained Exercise Time Max WorkLoad attained :9.2(Good Effort Tolerance) StageTime PhaseTime Speed 2:00 3:01 3:00 2:03 3:01 1:00 8:04 6:02 3:02 :08:03 :161 bpm 87% of Max Predictable HR 186 0.0 0.0 0.0 2.5 .7 Negatine Grade 14.0 12.0 0.0 0.0 0.0 0.0 10.0 9.2 7.1 1.0 1.0 4.7 METS 0 0 reor 107 123 140 136 161 109 H.R. 50 80 86 78 アスト 140/85 150/85 160/90 150/85 150/85 140/80 120/80 120/80 130/80 120/80 History 120/80 B.P. R.P.P. 210 241 149 66 90 4 03 96 96 93 60 ×100 PVC Comments Dr. Naresh Kumar Mohanka RMC No.: 35703

ABBS, DIP. CARDIO (ESCORTS) -0.3 PeakEx PreEx III Ξ 0.8 D.E.M. (RCGP-UK) avF **√**2 avR 46 5 **Y**4 చ ≾ avL = S 6 0.5 mm/Div 9 PR 12 15 18 21 Min.

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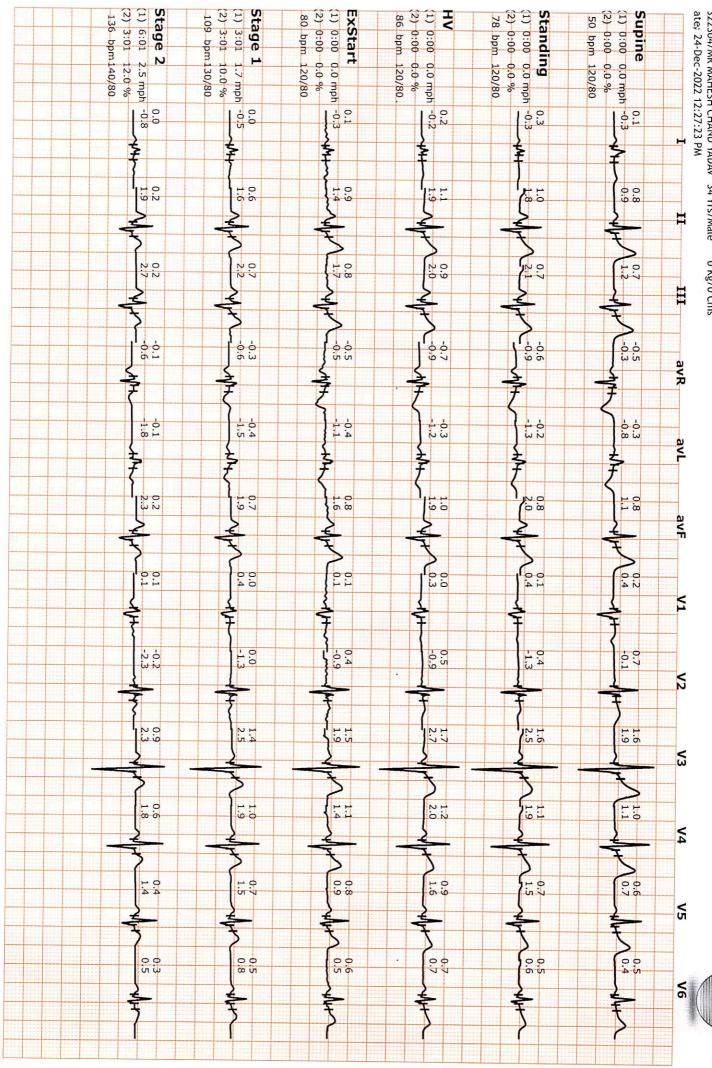
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17 Lead + Median

17 Lead + Median

1-14, Vidhyadhar Nagar Enclave, Phase -2, Jaipur 322304/MR MAHESH CHAND YADAV 34 Yrs/Male 0 Kg/0 Cms ate; 24-Dec-2022 12:27:23 PM



1-14, Vidhyadhar Nagar Enclave, Phase -2, Jaipur

322304/MR MAHESH CHAND YADAV 34 Yrs/Male 0 Kg/0 Cms

