



भारत सरकार

Government of India



भारती वर्मा

Bharti Verma

जन्म तिथि / DOB : 12/04/1992

पुरुष / Male



2004 6246 0164

मेरा आधार, मेरी पहचान



राष्ट्रीय विशिष्ट पहचान प्राधिकरण

Unique Identification Authority of India

पता:

वॉर्ड न 03, बलाईयो का मोहल्ला,
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~~2004-6246~~ 0164

मेरा आधार, मेरी पहचान

Bharti

Dr. PIYUSH GOYAL
MBBS, DMRD (Radiologist)
RMC No. 037041

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(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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General Physical Examination

Date of Examination: 13/01/2024

Name: Bharti Verma Age: 31 DOB: 12/04/1992 Sex: Female

Referred By: Bank of Baroda

Photo ID: Adhaar Card ID #: 0164

Ht: 157 (cm)

Wt: 66 (Kg)

Chest (Expiration): 95 (cm)

Abdomen Circumference: 87 (cm)

Blood Pressure: 120/80 mm Hg PR: 79 / min RR: 18 / min Temp: Afebrile

BMI 26.8

Eye Examination: R/E, 6/6, N/G, NCB

L/E 6/6 N/G NCB

Other: _____

On examination he/she appears physically and mentally fit: Yes / No

Signature Of Examinee: Bharti Name of Examinee: Bharti Verma

Signature Medical Examiner: Dr. PIYUSH GOYAL Name Medical Examiner: Dr. piyush goyal
MBBS, DMRD (Radiologist)
RMC No. 037041



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NAME :- Mrs. BHARTI VERMA

Age :- 31 Yrs 9 Mon 2 Days

Sex :- Female

Patient ID :-42234371

Date :- 13/01/2024 09:04:25

Ref. By Doctor:-BANK OF BARODA

Lab/Hosp :-

Company :- Mr.MEDIWHEEL

Final Authentication : 13/01/2024 17:41:01

HAEMOGARAM

HAEMATOTOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP BELOW 40 FEMAL			
HAEMOGLOBIN (Hb)	7.9 L	g/dL	12.0 - 15.0
TOTAL LEUCOCYTE COUNT	6.50	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	58.0	%	40.0 - 80.0
LYMPHOCYTE	35.0	%	20.0 - 40.0
EOSINOPHIL	3.0	%	1.0 - 6.0
MONOCYTE	4.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	4.39	$\times 10^6/\mu\text{L}$	3.80 - 4.80
HEMATOCRIT (HCT)	27.30 L	%	36.00 - 46.00
MEAN CORP VOLUME (MCV)	62.0 L	fL	83.0 - 101.0
MEAN CORP HB (MCH)	18.0 L	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	28.9 L	g/dL	31.5 - 34.5
PLATELET COUNT	413 H	$\times 10^3/\mu\text{L}$	150 - 410
RDW-CV	16.6 H	%	11.6 - 14.0



Tanu

DR.TANU RUNGTA

MD (Pathology)

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HAEMATOLOGY

Erythrocyte Sedimentation Rate (ESR)

24 H

mm in 1st hr

00 - 20

Method - Westergren

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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(CBC): Methodology: TLC,DLC Fluorescent Flow cytometry. HB-SLS method,TRBC,PCV,PLT Hydrodynamically focused Impedance, and MCH,MCV,MCHC,MENTZER INDEX are calculated. InstrumentName: Sysmex 6 part fully automatic analyzer XN-L,Japan





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BIOCHEMISTRY

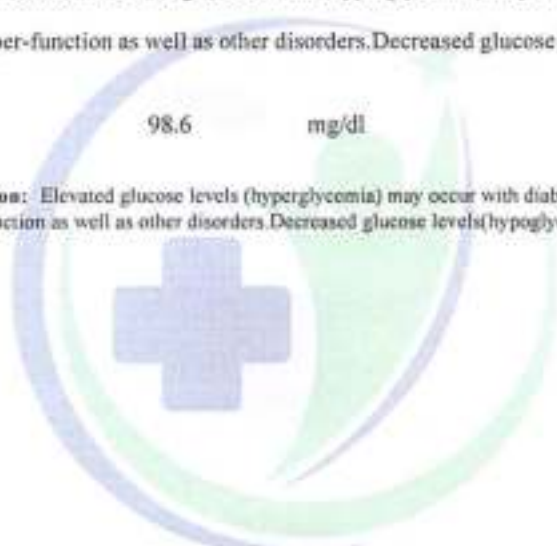
Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Method- GOD POD	81.8	mg/dl	70.0 - 115.0
Impaired glucose tolerance (IGT)	111 - 125 mg/dL		
Diabetes Mellitus (DM)	> 126 mg/dL		

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.

BLOOD SUGAR PP (Plasma)
Method- GOD PAP

98.6 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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HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
GLYCOSYLATED HEMOGLOBIN (HbA1C) Method - CAPILLARY with EDTA	5.1	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0
MEAN PLASMA GLUCOSE Method - Calculated Parameter	101	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: esp. vitamin B12 deficiency, decreased erythropoiesis

- Decreased HbA1c: administration of erythropoietin, esp. vitamin B12, reticulocytosis, chronic liver disease.

2. Altered Haemoglobin-Genetic or chemical alterations in hemoglobin, hemoglobinopathies, HbF, methemoglobin, may increase or decrease HbA1c.

3. Glycation

- Increased HbA1c: alcoholism, chronic renal failure, decreased intracellular 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 anorecticals, ribavirin & dapsone.

5. Others

- Increased HbA1c: hyperbilirubinemia, carbamylated hemoglobin, alcoholism, large doses of aspirin, chronic opiate use, chronic renal failure

- Decreased HbA1c: hyperglycemia, reticulocytosis, chronic liver disease, aspirin, vitamin C and E, splenomegaly, rheumatoid arthritis or drugs

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HAEMATOLOGY

BLOOD GROUP ABO

Method - Haemagglutination reaction

"A" POSITIVE



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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
TOTAL CHOLESTEROL Method- CHOD-PAP methodology	118.00	mg/dl	Desirable <200 Borderline 200-239 High > 240
<i>InstrumentName:MISPA PLUS Interpretation: Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.</i>			
TRIGLYCERIDES Method- GPO-PAP	70.90	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
<i>InstrumentName:Randox Rx Imola 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.</i>			
DIRECT HDL CHOLESTEROL Method- Direct clearance Method	43.80	mg/dl	MALE- 30-70 FEMALE - 30-85
<i>Instrument Name:Rx Daytona 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.</i>			
LDL CHOLESTEROL Method- Calculated Method	62.38	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method- Calculated	14.18	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method- Calculated	2.69		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method- Calculated	1.42		0.00 - 3.50
TOTAL LIPID Method- CALCULATED	356.20	L mg/dl	400.00 - 1000.00

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

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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 from peripheral tissues.



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BIOCHEMISTRY

LIVER PROFILE WITH GGT

SERUM BILIRUBIN (TOTAL)

Method- DMSO/Diaz

0.69 mg/dl.

Infants : 0.2-8.0 mg/dL
Adult - Up to - 1.2 mg/dL

SERUM BILIRUBIN (DIRECT)

Method- DMSO/Diaz

0.23 mg/dl.

Up to 0.40 mg/dL

SERUM BILIRUBIN (INDIRECT)

Method- Calculated

0.46 mg/dl

0.30-0.70

SGOT

Method- IFCC

20.5 U/L

0.0 - 40.0

SGPT

Method- IFCC

24.9 U/L

0.0 - 35.0

SERUM ALKALINE PHOSPHATASE

Method- DGKC - SCE

126.00 U/L

64.00 - 306.00

InstrumentName: MISPA PLUS **Interpretation:** Measurements of alkaline phosphatase are of use in the diagnosis, treatment and investigation of hepatobiliary disease and in bone disease associated with increased osteoblastic activity. Alkaline phosphatase is also used in the diagnosis of parathyroid and intestinal disease.

SERUM GAMMA GT

Method- Sraz methodology

Instrument Name: Randox Rr level

Interpretation: Elevations in GGT levels occur earlier and more pronounced than those with other liver enzymes in cases of obstructive jaundice and

metastatic neoplasms. It may reach 5 to 30 times normal levels in severe or post

hepatic biliary obstruction. Only moderate elevations in the enzyme level (2 to 3 times normal) are observed with infectious hepatitis.

21.60 U/L

5.00 - 32.00

SERUM TOTAL PROTEIN

Method- Direct Biuret Reagent

6.89 g/dl

6.00 - 8.40

SERUM ALBUMIN

Method- Bromocresol Green

3.97 g/dl

3.50 - 5.50

SERUM GLOBULIN

Method- CALCULATION

2.92 gm/dl

2.20 - 3.50

A/G RATIO

1.36

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.

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 cases) on those individuals taking certain medications, such as

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RFT / KFT WITH ELECTROLYTES

SERUM UREA Method- Urnase/GLDH	21.50	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 Method- Jaffe's Method	0.91	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	4.58	mg/dl	2.40 - 7.00
-----------------	------	-------	-------------

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

SODIUM Method- ISE	136.7	mmol/L	135.0 - 150.0
-----------------------	-------	--------	---------------

POTASSIUM Method- ISE	3.67	mmol/L	3.50 - 5.50
--------------------------	------	--------	-------------

CHLORIDE Method- ISE	98.9	mmol/L	94.0 - 110.0
-------------------------	------	--------	--------------

SERUM CALCIUM Method- Arsmazo III Method	9.68	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 Method- Direct Buret Reagent	6.89	g/dl	6.00 - 8.40
-----------------------------------------------------	------	------	-------------

SERUM ALBUMIN Method- Bromocresol Green	3.97	g/dl	3.50 - 5.50
--------------------------------------------	------	------	-------------

SERUM GLOBULIN Method- CALCULATION	2.92	gm/dl	2.20 - 3.50
---------------------------------------	------	-------	-------------

A/G RATIO	1.36		1.30 - 2.50
-----------	------	--	-------------

Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of dis... liver, kidney and

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BIOCHEMISTRY

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 blood increases. 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.

Apart from renal failure Blood Urea can increase in dehydration and GI bleed.



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CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
Urine Routine			
PHYSICAL EXAMINATION			
COLOUR	PALE YELLOW		PALE YELLOW
APPEARANCE	Clear		Clear
CHEMICAL EXAMINATION			
REACTION(PH)	5.5		5.0 - 7.5
SPECIFIC GRAVITY	1.015		1.010 - 1.030
PROTEIN	NIL		NIL
SUGAR	NIL		NIL
BILIRUBIN	NEGATIVE		NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVE		NEGATIVE
NITRITE	NEGATIVE		NEGATIVE
MICROSCOPY EXAMINATION			
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		ABSENT



Tanu

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

Technologist
VIKARANTSI
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(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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Central Spine, Vidhyadhar Nagar, Jaipur - 302023
📞 +91 141 4824885 📧 maxcarediagnostics1@gmail.com



NAME :- Mrs. BHARTI VERMA	Patient ID :-12234371	Date :- 13/01/2024	09:04:25
Age :- 31 Yrs 9 Mon 2 Days	Ref. By Doctor:-BANK OF BARODA		
Sex :- Female	Lab/Hosp :-		
	Company :- Mr.MEDIWHEEL		

Final Authentication : 13/01/2024 17:41:01

CLINICAL PATHOLOGY

STOOL ANALYSIS

PHYSICAL EXAMINATION

MUCUS

BLOOD

MICROSCOPIC EXAMINATION

RBC's

/HPF

WBC/HPF

/HPF

OVA

CYSTS

OTHERS

Collected Sample Received



Technologist
VIKARANJJI
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TOTAL THYROID PROFILE

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
THYROID-TRIiodothyronine T3 Method- ECLIA	1.20	ng/ml.	0.70 - 2.04 0.70-2.04

NOTE: In pregnancy total T3,T4 increase to 1.5 times the normal range.

Reference Range (T3): Premature infants 26-30 Weeks ,3-4 days

Full-Term infants 1-3 days

1 Week

1- 11 Months

Prepubertal Children

0.24 - 1.32 ng/ml

0.89 - 4.05 ng/ml

0.91 - 3.00 ng/ml

0.85 - 2.50 ng/ml

1.19 - 2.18 ng/ml

Reference Ranges (T4): Premature infants 26-30 weeks ,3-4 days

Full -Term infants 1-3 days

1 weeks 6.00 - 15.9 ug/dl 1-11 Months

Prepubertal children 12 months-2yrs

Prepubertal children 3-9 yrs

2.60 - 14.0 ug/dl

8.20 - 19.9 ug/dl

6.10 - 14.9 ug/dl

6.80 - 13.5 ug/dl

5.50 - 12.8 ug/dl

Reference Ranges (TSH): Premature Infants 26-32 weeks ,3-4 Days

Full Term infants 4 Days

0.80 - 6.9 uIU/ml

1.36 - 16 uIU/ml

1 - 11 Months:0.80 - 7.70 | Prepubertal children:0.60 - 5.50. Primary malfunction of the thyroid gland may result in hyper or low release of T3 or T4

In addition as TSH directly affect thyroid function malfunction of the pituitary or the hypothalamus influences the thyroid gland activity. Disease in any portion of the thyroid pituitary hypothalamus system may influence the level of T3 and T4 in the blood in Primary hypo thyroidism TSH levels

in secondary and tertiary hypothyroidism TSH levels may be low

THYROID-THYROXINE (T4)
Method- ECLIA

5.10 - 14.10

5.1-14.1

NOTE-TSH levels are subject to circadian variation,peaking 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 measured 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.simultaneous measurement of TSH with free T4 is useful in evaluating differential diagnosis.

INTERPRETATION-Ultra Sensitive 4th generation assay 1.Primary hyperthyroidism is accompanied by raised 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 antibody increased seen in patients with Hashimoto's thyroiditis 5.HighTSH,low FT4 and Thyroid microsomal antibody normal seen in patients with iodine deficiency/Congenital T4 synthesis deficiency 6.Low TSH,Low FT4 and TSH 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 Thyrotoxicosis 9.Normal or T3 & T4 10.Normal T3 & T4 along with TSH indicate mild / Subclinical Hyperthyroidism. 11.Normal T3 & T4 along with TSH is seen in Hypothyroidism. 12.Normal T3 & T4 levels 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 catabolism 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 radioactive stain 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 concentration with age, and it is debatable whether this is due to a real change with age or an increasing proportion of unmeasured thyroid disease in the elder.

TSH

Method- ECLIA

4.437

uIU/ml.

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IMMUNOASSAY

4th Generation Assay,Reference ranges vary between laboratories

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.

NOTE-TSH levels are subject to circadian 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.

INTERPRETATION

- 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with ↓ TSH level.
- 2.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & ↑serum TSH levels
- 3.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis
- 4.Normal or ↓ T3 & ↑T4 levels indicate T4 Thyrotoxicosis (problem is conversion of T4 to T3)
- 5.Normal T3 & T4 along with ↓ TSH indicate mild / Subclinical Hyperthyroidism

COMMENTS: 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.

Disclaimer-TSH is an important marker for the diagnosis of thyroid dysfunction. Recent studies have shown that the TSH distribution progressively shifts to a higher concentration with age, and it is debatable whether this is due to a real change with age or an increasing proportion of unrecognized thyroid disease in the elderly.

Reference ranges are from Teltz fundamental of clinical chemistry 8th ed (2018)

Test performed by Instrument : Beckman coulter Dxi 800

Note : The result obtained relate only to the sample given/ received & tested. A single test result is not always indicative of a disease, it has to be correlated with clinical data for interpretation.

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PAP SMEAR

PAP SMEAR FOR CYTOLOGY EXAMINATION

Microscopic & diagnosis,

Smears are

No endocervical cells seen.

No atypical or malignant cells seen.

IMPRESSION :Inflammatory smears, (Negative for intraepithelial lesion or malignancy).

Adv: Clinical correlation.

Note: Please note papanicolaou smear study is a screening procedure for cervical cancer with inherent false negative result, hence should be interpreted with caution.

*** End of Report ***

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MRS. BHARTI VERMA	Age : 31 Y/F
Registration Date: 13/01/2024	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (149 mm). Echo-texture is normal. No focal space occupying lesion is seen within liver parenchyma. Intrahepatic biliary channels are not dilated. Portal vein diameter is normal.

Gall bladder is well distended. Wall is not thickened. 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. Echotexture is normal. No focal lesion is seen.

Kidneys are normally sited and are of normal size and shape. Cortico-medullary echoes are normal. No focal lesion is seen. Collecting system does not show any dilatation or calculus.

Right kidney is measuring approx. 107 mm.

Left kidney is measuring approx. 102 mm.

Urinary bladder is sub-optimally distended and does not show any calculus or mass lesion.

Uterus is anteverted and retroflexed (measuring approx. 83 x 52 mm).

Myometrium shows normal echo -pattern. No focal space occupying lesion is seen. Endometrial echo is normal. Endometrial thickness is 7.4 mm.

Both ovaries are visualized and are normal. No adnexal mass lesion is seen.

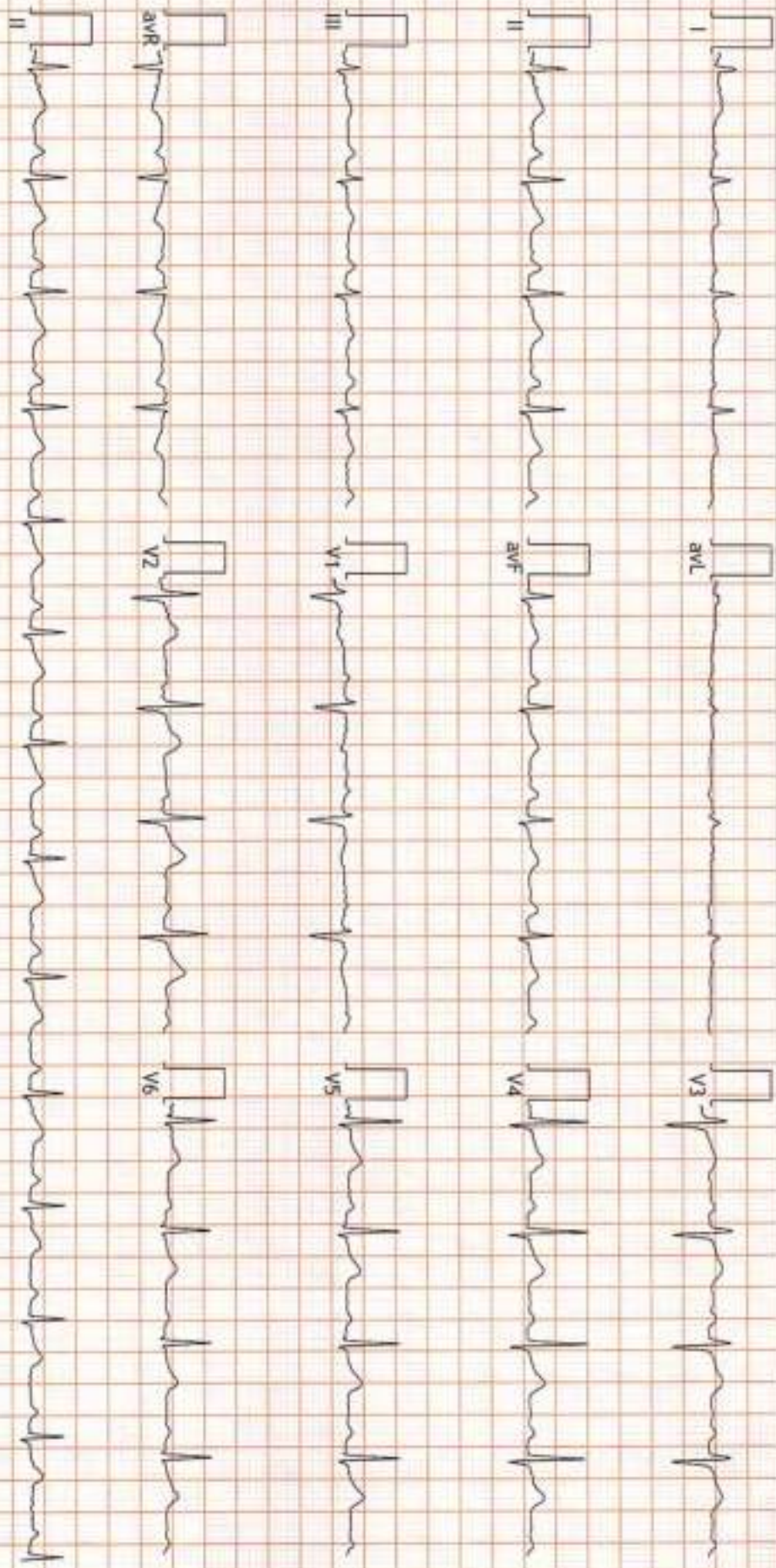
No enlarged nodes are visualized. No retro-peritoneal lesion is identified.

No significant free fluid is seen in pouch of Douglas.

IMPRESSION: No significant abnormality is detected.

Dr. Mukesh Sharma
M.B.B.S; M.D. (Radiodiagnosis)
RMC No. 43418/17437

Dr. MUKESH SHARMA
M.B.B.S., M.D.(Radiodiagnosis)
RMC No. : 43418/17437
P3 Health Solutions LLP



FINDINGS: Normal Sinus Rhythm
Vent Rate : 79 bpm; PR Interval : 180 ms; QRS Duration: 90 ms; QT/QTc Int : 374/430 ms
P-QRS-T axis: 72 • 36 • 53 • (Deg)
Comments :

TWML

Bharti

MS

Dr. Naresh Kumar Mohanka
RMC No: 35703
MBBS, Dip in Cardiology (ESCORTS)
D.E.M. (RCGP-UK)

For the safety of your staff, please do not
allow them to work in this area if you
are not with them.
Any job staff leaves the site
during duty hours and the camera
is not in the region, their
activity will be
monitored.

This is believed to
have a staff member.



GPS Map Camera

Jaipur, Rajasthan, India

Vidhyadhar Enclave II, b 14, Sector 2 Rd, Sector 2, Central Spine, Vidyadhar
Nagar, Jaipur, Rajasthan 302039, India

Lat 26.964455°

Long 75.782581°

13/01/24 01:04 PM GMT +05:30

Google



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NAME:	MRS. BHARTI VERMA	AGE	31 YRS/F
REF.BY	BANK OF BARODA	DATE	13/01/2024

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

Dr. Mukesh Sharma
M.B.B.S; M.D. (Radiodiagnosis)
RMC No. 43418/17437



 GPS Map Camera

Jaipur, Rajasthan, India

Vidhyadhar Enclave II, b 14, Sector 2 Rd, Sector 2, Central Spine, Vidyadhar Nagar, Jaipur, Rajasthan 302039, India

Lat 26.964515°

Long 75.782521°

13/01/24 01:06 PM GMT +05:30



R



12234371 BHARTI VERMA 31 YRS MEDIWHEEL F

13 JAN 2024

MAXCARE DIAGNOSTIC (ASSOCIATES OF P3 HEALTH SOLUTIONS LLP)

