



## List of Pending Test

<u>Test Name</u>	<u>Status</u>
ESR	Ready
UR (URINE ROUTINE)	Ready
Lipid Profile	Ready
BUN	Ready
Uric Acid	Ready
Calcium	Ready
Bilirubin	Ready
SGOT	Ready
SGPT	Ready
Alk. Phosphatase	Ready
GGTP	Ready
Proteins	Ready
TSH	Ready
CREATININE	Ready
CBC	Ready
VITAMIN B12	Ready
HbA1C	Ready
IRON & TIBC	Ready
VITAMIN D	Ready
BSF	Ready

**Note:-** Following department reports if any,hardcopy of the reports to be collected from the centre where test is performed.

- E.C.G
  - STRESS TEST
  - PFT
  - BMD
  - EEG/EMG/NCV
- and some special pathological test.



# P. H. DIAGNOSTIC CENTRE

ISO 9001 : 2015 Certified

**PATIENT'S NAME** : MRS. JAIN PAPITA  
**AGE / GENDER** : 43 Years / Female  
**REFERRED BY DR** : KERING RAMESH  
**PATIENT ID** : 701005437  
**SAMPLE COLLECTED BY** : P.H.AUNDH



**CLIENT** : PH COM  
**REGISTRATION DATE** : 24/06/2021 08:57am  
**SAMPLE COLL. DATE** : 24/06/2021 09:06am  
**ACCESSION DATE** : 24/06/2021 12:00pm  
**AUTHENTICATION DATE** : 24/06/2021 01:26pm

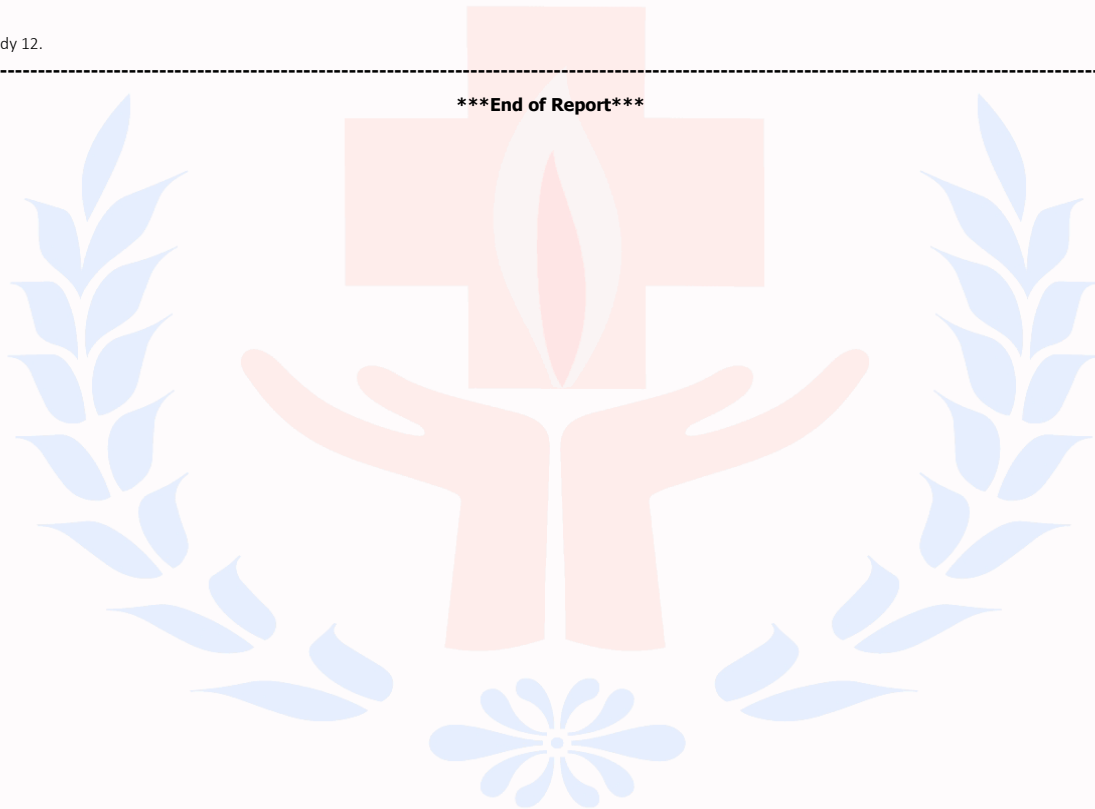
## ERYTHROCYTE SEDIMENTATION RATE

<u>Test</u>	<u>Observed Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
ESR	07	mm/hr	0 - 12

Specimen : Whole Blood  
Method : Modified Westergren

Instrument : Sedy 12.

\*\*\*End of Report\*\*\*



MC - 2630

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## COMPLETE BLOOD COUNT

Test	Observed Value	Unit	Biological Reference Interval
HAEMOGLOBIN	12.0	g/dL	12.0 - 15.0
R.B.C COUNT	3.83	10 <sup>6</sup> / uL	3.8 - 4.8
PCV	<b>L 34.7</b>	%	36 - 46
MCV	90.60	fL	83 - 101
MCH	31.33	pg	27 - 32
MCHC	<b>H 34.58</b>	g/dl	31.5 - 34.5
RDW	12.9	%	11.0 - 14.5
PLATELET COUNT	241	x 10 <sup>3</sup> /uL	150 - 410
MEAN PLATELET VOLUME(MPV)	8.1	fL	7.8 - 11.0
W.B.C COUNT	5100	per cu-mm	4000 - 10000

## DIFFERENTIAL COUNT

NEUTROPHILS	58.2	%	40.0 - 75.0
LYMPHOCYTES	30.9	%	20 - 45
EOSINOPHILS	1.8	%	1.0 - 6.0
MONOCYTES	8.2	%	0.0 - 10.0
BASOPHILS	0.9	%	0.0 - 1.0
ABSOLUTE NEUTROPHIL COUNT	2968	per cumm	2000 - 7000
ABSOLUTE LYMPHOCYTE COUNT	1576	per cumm	1000 - 3000
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.9		< 3
RBC MORPHOLOGY	Normocytic Normochromic		
W.B.C MORPHOLOGY	Normal		
PLATELET MORPHOLOGY	Platelet adequate		

Specimen : Whole Blood (EDTA)

Method : Coulter Principle/Derived from WBC Histogram/Cyanmethaemoglobin photometry/Calculated.

Instrument : Beckmen Coulter LH750/DXH800/Microscopy.

\*\*\*End of Report\*\*\*



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## BIOCHEMICAL TEST

Test	Observed Value	Unit	Biological Reference Interval
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<b>CALCIUM</b>	8.9	mg/dL	8.6 - 10.0
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Serum by NM-BAPTA.

Instrument : Cobas 6000 / Cobas C 311

### **Bilirubin**

<b>BILIRUBIN (TOTAL)</b>	0.38	mg/dL	0.3 - 1.2
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<b>BILIRUBIN (DIRECT)</b>	0.14	mg/dL	0 - 0.4
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<b>BILIRUBIN (INDIRECT)</b>	0.24	mg/dL	0.1 - 1.0
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Serum by Diazo Method

Instrument : Cobas 6000 / Cobas C 311

<b>SGOT (AST)</b>	15.7	U/L	0 - 32
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Serum by IFCC Method, Kinetic

Instrument : Cobas 6000 / Cobas C 311

<b>SGPT (ALT)</b>	11.0	U/L	0 - 33
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Serum by IFCC Method, Kinetic

Instrument : Cobas 6000 / Cobas C 311

<b>ALKALINE PHOSPHATASE</b>	45	U/L	35 - 105
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Serum by IFCC Method

Instrument : Cobas 6000 / Cobas C 311

<b>(GGT) GAMMA- GLUTAMYLTRANSFERASE</b>	13.3	U/L	< 40
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Serum by IFCC / SZASZ

Instrument : Cobas 6000 / Cobas C 311

<b>BLOOD UREA</b>	14.6	mg/dL	12.8 - 42.8
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<b>BLOOD UREA NITROGEN</b>	6.82	mg/dL	6 - 20
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Serum by Urease - GLDH Method

Instrument : Cobas 6000/ Cobas C311

<b>URIC ACID</b>	5.1	mg/dL	2.4 - 5.7
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Serum by Enzymatic colorimetric test (Uricase)

Instrument : Cobas 6000 / Cobas C 311



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## BIOCHEMICAL TEST

Test	Observed Value	Unit	Biological Reference Interval
<b>TOTAL PROTEIN</b>	6.56	g/dL	6.4 - 8.3
<b>ALBUMIN</b>	4.33	g/dL	3.5 - 5.2
<b>GLOBULIN</b>	2.23	g/dL	2.3 - 3.5
<b>A/G RATIO</b>	1.94		0.9 - 2.0

Serum by Biuret Method (Protein), BCG Gen 2 (Albumin)  
Instrument : Cobas 6000 / Cobas C 311

### SERUM CREATININE

0.70

mg/dL

0.5 - 0.9

Serum by Jaffe method Gen 2  
Instrument : Cobas 6000 / Cobas C 311

\*\*\*End of Report\*\*\*



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## LIPID PROFILE

Test	Observed Value	Unit	Biological Reference Interval
<b>BIO-CHEMICAL TEST</b>			
<b>TOTAL CHOLESTEROL</b>	<b>H 227.8</b>	mg/dL	< 200
<b>TRIGLYCERIDES</b>	136.6	mg/dL	0 - 150
<b>HDL CHOLESTEROL</b>	43.8	mg/dL	30 - 70
<b>VLDL CHOLESTEROL</b>	27.32	mg/dL	Upto 35
<b>LDL CHOLESTEROL</b>	<b>H 156.68</b>	mg/dL	< 100
<b>TC/HDLC RATIO</b>	<b>H 5.20</b>		Upto 5.0
<b>LDLC/HDLC RATIO</b>	3.58		2.5 - 3.5

Specimen : Serum

Method : CHOD-POD (cholesterol/Enzymatic colorimetric (triglyceride, LDL)/PEG(HDL)/Calculated(VLDL)

Instrument Used : Cobas 6000/Cobas c 311.

INTERPRETATION : As per NCEP 2001 ATP III guidelines May 2001

Total Cholesterol

200 - 239 : Borderline high

> 240 : High

Triglyceride

150 - 199 : Borderline high

200 - 499 : High

> 500 : Very high

LDL Cholesterol

100 - 129 : Near / Above optimal

130 - 159 : Borderline high

160 - 189 : High

>190 : Very High

\*\*\*End of Report\*\*\*



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## S.IRON & TIBC

<u>Test</u>	<u>Observed Value</u>	<u>Unit</u>	<u>Biological Reference Interval</u>
<b>IRON</b>	82.78	µg/dL	33 - 193
<b>TOTAL IRON BINDING CAPACITY (TIBC)</b>	339	µg/dL	255 - 450
<b>IRON SATURATION</b>	24.42	%	20 - 50

Serum by Ferrozine Method - without deproteinization (Iron).  
Direct Determination (TIBC).  
Calculation (Saturation).

Instrument : Cobas 6000 / Cobas C 311.

### INTERPRETATION :

1. In Iron deficiency anemia the TIBC is elevated and the Iron saturation is lowered to 15% or less.
2. Low serum Iron associated with low TIBC is characteristic of anemia of chronic disorders, malignant tumors and infections.
3. Estrogens and oral contraceptives increase TIBC levels.

\*\*\*End of Report\*\*\*



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## URINE ROUTINE

Test	Observed Value	Unit	Biological Reference Interval
<b><u>PHYSICAL EXAMINATION</u></b>			
QUANTITY	20 ml		
COLOUR	Pale yellow		Pale yellow
APPEARANCE	Clear		Clear
PH	6.5		4.5- 8
SPECIFIC GRAVITY	1.015		1.005 - 1.030
DEPOSIT	Absent		Absent
<b><u>CHEMICAL EXAMINATION</u></b>			
PROTEINS	Negative		Negative
GLUCOSE	Negative		Negative
KETONE BODIES	Negative		Negative
OCCULT BLOOD	Positive		Negative
BILIRUBIN	Negative		Negative
UROBILINOGEN	Normal		Normal
NITRITE	Negative		Negative
<b><u>MICROSCOPIC EXAMINATION :-</u></b>			
PUS CELLS	Occasional		0-2 / hpf
RED BLOOD CELLS	3 - 4/hpf		0-2 / hpf
EPITHELIAL CELLS	2 - 3 /hpf		< 20 / hpf

Method :

Visual observation/GOD-POD (Glucose)/Protein error (Protein)/Legal's test (Ketone)/Peroxidase (Occult blood)/PH indicator (PH)/Diazonium Coupling (Bilirubin)/Diazonium reaction (urobilinogen)/Griess test (Nitrite)/Ionic concentration (Specific Gravity)/wet mount/Microscopy.

Instrument : Cobas u 411.

\*\*\*End of Report\*\*\*



MC - 2630

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Reviewed By : MANISHA AMBLE





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## BLOOD GLUCOSE FASTING

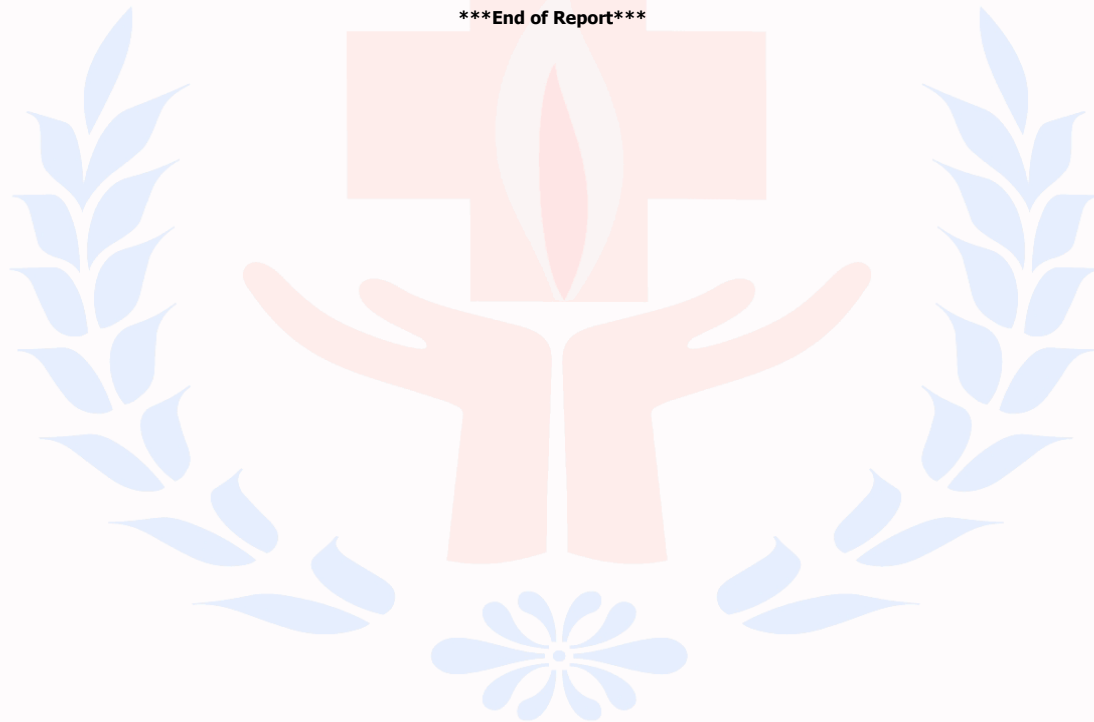
<u>Test</u>	<u>Observed Value</u>	<u>Unit</u>	<u>Biological Reference Interval</u>
<b>BLOOD GLUCOSE FASTING</b>	89.0	mg/dL	70 -110

Specimen :Plasma.

Method : Hexokinase (UV).

Instrument : Cobas 6000 / Cobas c 311.

\*\*\*End of Report\*\*\*



MC - 2630

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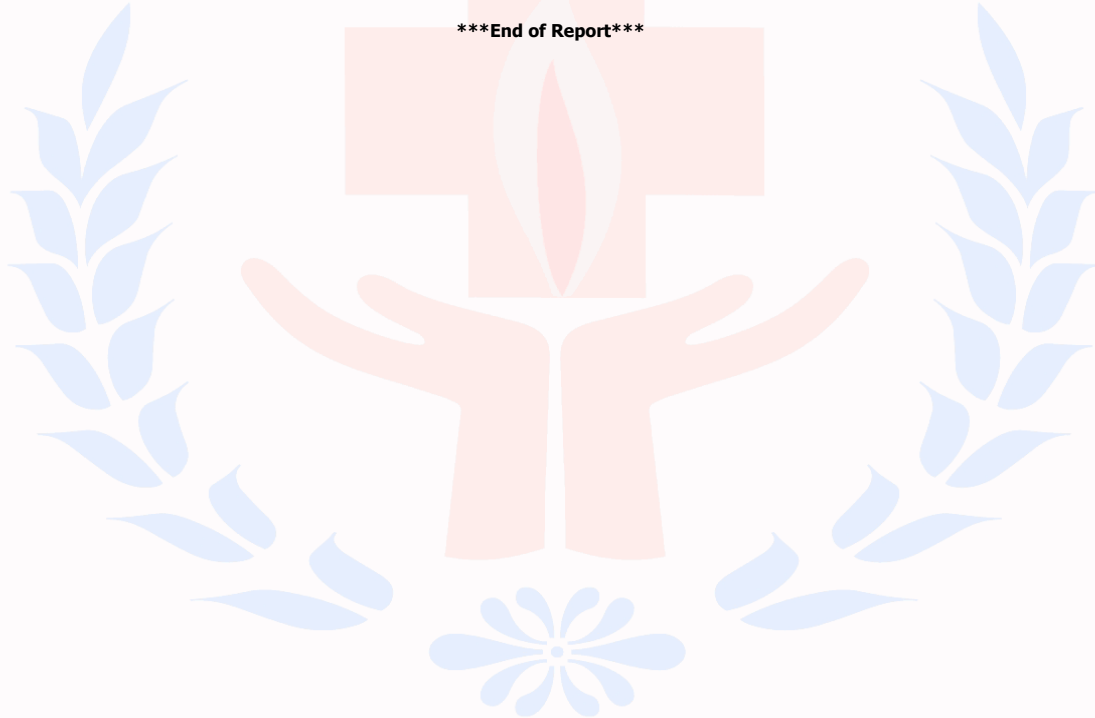
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## TSH

<u>Test</u>	<u>Observed Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
<b>THYROID STIMULATING HORMONE (TSH)</b>	<b>L 0.221</b>	µIU/mL	0.4 - 4.5 1st Trimester : 0.3 - 4.5 2nd Trimester : 0.5 - 4.6 3rd Trimester : 0.8 - 5.2

Specimen : Serum By CMIA  
Instrument: ARCHITECT i2000 SRPLUS

\*\*\*End of Report\*\*\*



MC - 2630

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## VITAMIN B12

<u>Test</u>	<u>Observed Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
<b>VITAMIN B12</b>	541.3	pg/mL	Normal : 180 - 914 Indeterminate : 145 - 180 Deficient : < 145

Specimen: Serum By ECLIA  
Instrument: Cobas 6000

### INTERPRETATION

1. Increased level are seen in Chronic granylocytic leukemia, COPD, Chronic renal leukocytosis, Liver cell damage, Obesity, Polycythemia vera, Severe CHF and some carcinomas.
2. Decreased level are seen in Abnormalities of cobalamin transport or metabolism, Bacterial overgrowth, Dietary deficiency, Gastric or small intestine surgery, Inflammatory bowel disease, Intestinal malabsorption, Intrinsic factor deficiency and Late pregnancy.
3. Pregnany, smoking, hemodialysis, multiple myeloma, can decrease B 12 levels.
4. Patients taking vitamin B12 supplementation may have misleading results.
5. A normal serum B12 level does not rule out tissue deficiency of vitamin B12.

\*\*\*End of Report\*\*\*



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## HbA1C

Test	Observed Value	Unit	Biological Reference Interval
HbA1C	5.5	%	Normal : 4.0 - 5.6 Pre Diabetes : 5.7 - 6.4 Diabetic : > 6.5
MEAN GLUCOSE LEVEL	111.15	mg/dL	

Specimen : Whole Blood EDTA  
Method : HPLC

### INTERPRETATION :

ADA Recommendation for Diabetic control

- 4 - 6 : Non-diabetic
- 6 - 7 : Excellent Control
- 7 - 8 : Fair To Good Control
- 8 - 10 : Unsatisfactory Control
- Above 10 : Poor Control

- HbA1c is used for monitoring diabetic control and reflects mean plasma glucose over three months.
- HbA1c is falsely low in diabetic with hemolytic disease. In these individuals a plasma fructosamine level may be used which evaluates diabetes over 15 days.
- Trends in HbA1c are a better indicator of diabetic control than a solitary test.
- HbA1c value is used to estimate the mean plasma Glucose(MPG) level over the last 90 days.

\*\*\*End of Report\*\*\*



MC - 2630

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## VITAMIN D TOTAL (25 HYDROXY )

<u>Test</u>	<u>Observed Value</u>	<u>Unit</u>	<u>Biological Reference Interval</u>
<b>VITAMIN D TOTAL (25-HYDROXY VIT.D)</b>	55.4	ng/mL	Deficiency : < 10 Insufficiency : 10 - 30 Sufficiency : 30 - 100 Toxicity : > 100

**METHOD** CMIA

### INTERPRETATION:

- Decreased in Malabsorption, Steatorrhea, Dietary osteomalacia, anticonvulsant osteomalacia, Biliary & portal cirrhosis, Thyrotoxicosis, Pancreatic insufficiency, Celiac disease, Inflammatory bowel disease, Rickets, Alzheimer disease.
- Increased in Vitamin D intoxication, Excessive exposure to sunlight.

\*\*\*End of Report\*\*\*



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