

Visit ID	: YGT41532	UHID/MR No	: YGT.0000041384
Patient Name	: Mrs. BANDELA RAJYA LAKSHMI	Client Code	: 1409
Age/Gender	: 27 Y 4 M 14 D /F	Barcode No	: 10816264
DOB	: 12/Jul/1996	Registration	: 25/Nov/2023 09:14AM
Ref Doctor	: SELF	Collected	: 25/Nov/2023 09:14AM
Client Name	: MEDI WHEELS	Received	:
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 25/Nov/2023 11:02AM
Hospital Name	:		

### **ULTRASOUND WHOLE ABDOMEN & PELVIS**

Clinical Details : General check-up.

**LIVER :** Normal in size (13.0 cm) and echo-texture. Intra hepatic biliary channels are not dilated. Visualised common bile duct & portal vein appears normal. **A 12 x 12 mm well** *defined hyperechoic lesion noted in segment VI of right lobe of liver - Likely Hemangioma.* 

GALL BLADDER : Well distended. No evidence of calculi / wall thickening.

**PANCREAS :** Normal in size and outlines. Parenchymal texture normal. No ductal dilatation. No calcifications / calculi.

SPLEEN : Normal in size (9.9 cm) and echotexture. No focal lesion is seen.

**RIGHT KIDNEY :** measures 10.8 x 4.5 cm. Normal in size with smooth contours. Parenchymal texture normal. No focal lesion is seen. Cortico-medullary differentiation well maintained. Collecting system does not show any dilatation or calculus.

**LEFT KIDNEY :** measures 10.9 x 4.4 cm. Normal in size with smooth contours. Parenchymal texture normal. No focal lesion is seen. Cortico-medullary differentiation well maintained. Collecting system does not show any dilatation. *3 mm calculus noted in upper pole of left kidney.* 

**URINARY BLADDER :** Well distended. No evidence of calculi or wall thickening.

**UTERUS :** Anteverted, measures 9.2 x 4.4 x 5.3 cm, normal in size. Myometrium shows normal echo-texture. No focal lesion is seen. *Endometrium is thickened and measures 13 mm.* 

Right ovary measures 3.2 x 2.7 cm and left ovary measures 2.8 x 2.2 cm. Both ovaries are normal in size & echotexture. No adnexal lesion seen.

No enlarged nodes are visualised. No retro-peritoneal lesion is identified. Great vessels appear normal.

No free fluid is seen in pelvis.

Verified By :



zustrmat.

Dr.SUSHMA VUYYURU MBBS;MD(Radio-Diagnosis) CONSULTANT RADIOLOGIST





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## IMPRESSION:

- Well defined hyperechoic lesion in segment VI of right lobe of liver Likely Hemangioma.
- Left renal calculus.
- Thickened endometrium.

Verified By :



zustrmar.

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## **X-RAY CHEST PA VIEW**

### Findings:

Soft tissues/ bony cage normal.

Trachea and Mediastinal structures are normal.

Heart size and configuration are normal.

Aorta and pulmonary vascularity are normal.

Lung parenchyma and CP angles are clear.

Bilateral hilae and diaphragmatic contours are normal.

# IMPRESSION :

• No Significant Abnormality Detected.

Suggested Clinical Correlation & Follow up.

Verified By : GOPI



Lushmar.

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DEPARTMENT OF HAEMATOLOGY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

ESR (ERYTHROCYTE SEDIMENTATION RATE)							
Sample Type : WHOLE BLOOD EDTA							
ERYTHROCYTE SEDIMENTATION RATE	70	mm/1st hr	0 - 15	Capillary Photometry			
COMMENTS: ESR is an acute phase reactant which indicates presence and intensity of an inflammatory process. It is never diagnostic of a specific disease. It is used to monitor the course or response to treatment of certain diseases. Extremely high levels are found in cases of malignancy, hematologic diseases, collagen disorders and renal diseases.							

Increased levels may indicate: Chronic renal failure (e.g., nephritis, nephrosis), malignant diseases (e.g., multiple myeloma, Hodgkin disease, advanced Carcinomas), bacterial infections (e.g., abdominal infections, acute pelvic inflammatory disease, syphilis, pneumonia), inflammatory diseases (e.g. temporal arteritis, polymyalgia rheumatic, rheumatoid arthritis, rheumatic fever, systemic lupus erythematosus [SLE]), necrotic diseases (e.g., acute myocardial infarction, necrotic tumor, gangrene of an extremity), diseases associated with increased proteins (e.g., hyperfibrinogenemia, macroglobulinemia), and severe anemias (e.g., iron deficiency or B12 deficiency).

Falsely decreased levels may indicate: Sickle cell anemia, spherocytosis, hypofibrinogenemia, or polycythemia vera.

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DEPARTMENT OF HAEMATOLOGY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

BLOOD GROUP ABO & RH Typing						
Sample Type : WHOLE BLOOD EDTA						
ABO	А					
Rh Typing	POSITIVE					
Method : Hemagglutination Tube method by forward and reverse grouping						
COMMENTS:						
The test will detect common blood are	uning system A B O	AB and Phosus (E	D) Unusual blood	arouns or rare subtypes		

The test will detect common blood grouping system A, B, O, AB and Rhesus (RhD). Unusual blood groups or rare subtypes will not be detected by this method. Further investigation by a blood transfusion laboratory, will be necessary to identify such groups.

**Disclaimer:** There is no trackable record of previous ABO & RH test for this patient in this lab. Please correlate with previous blood group findings. Advsied cross matching before transfusion

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DEPARTMENT OF HAEMATOLOGY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

CBC	C(COMPLE	TE BLOOD CO	DUNT)	
Sample Type : WHOLE BLOOD EDTA				
HAEMOGLOBIN (HB)	12.7	g/dl	12.0 - 15.0	Cyanide-free SLS method
RBC COUNT(RED BLOOD CELL COUNT)	4.40	million/cmm	3.80 - 4.80	Impedance
PCV/HAEMATOCRIT	37.2	%	36.0 - 46.0	RBC pulse height detection
MCV	84.5	fL	83 - 101	Automated/Calculated
МСН	28.7	pg	27 - 32	Automated/Calculated
MCHC	34.0	g/dl	31.5 - 34.5	Automated/Calculated
RDW - CV	12.6	%	11.0-16.0	Automated Calculated
RDW - SD	41.3	fl	35.0-56.0	Calculated
MPV	7.8	fL	6.5 - 10.0	Calculated
PDW	15.5	fL	8.30-25.00	Calculated
PCT	0.33	%	0.15-0.62	Calculated
TOTAL LEUCOCYTE COUNT	8,910	cells/ml	4000 - 11000	Flow Cytometry
DLC (by Flow cytometry/Microscopy)				
NEUTROPHIL	60	%	40 - 80	Impedance
LYMPHOCYTE	32	%	20 - 40	Impedance
EOSINOPHIL	03	%	01 - 06	Impedance
MONOCYTE	05	%	02 - 10	Impedance
BASOPHIL	0	%	0 - 1	Impedance
PLATELET COUNT	4.18	Lakhs/cumm	1.50 - 4.10	Impedance



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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

THYROID PROFILE (T3,T4,TSH)							
Sample Type : SERUM							
T3	0.93	ng/ml	0.60 - 1.78	CLIA			
T4	6.63	ug/dl	4.82-15.65	CLIA			
TSH	4.70	ulU/mL	0.30 - 5.60	CLIA			
				·			

#### INTERPRETATION:

1. Serum T3, T4 and TSH are the measurements form three components of thyroid screening panel and are useful in diagnosing various disorders of thyroid gland function.

2. Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values along with depressed TSH levels.

3. Primary hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels.

4. Normal T4 levels accompanied by high T3 levels are seen in patients with T3 thyrotoxicosis. Slightly elevated T3 levels may be found in pregnancy and in estrogen therapy while depressed levels may be encountered in severe illness, mainutrition, renal failure and during therapy with drugs like propanolol and propylthiouracil.

5. Although elevated TSH levels are nearly always indicative of primary hypothyroidism, rarely they can result from TSH secreting pituitary tumors (secondary hyperthyroidism)

6. Low levels of Thyroid hormones (T3, T4 & FT3, FT4) are seen in cases of primary, secondary and tertiary hypothyroidism and sometimes 7. Increased levels are found in Grave's disease, hyperthyroidism and thyroid hormone resistance.

8. TSH levels are raised in primary hypothyroidism and are low in hyperthyroidism and secondary hypothyroidism.

9.	REFERENCE RANGE :				
	PREGNANCY	TSH in ul U/mL			
	1st Trimester	0.60 - 3.40			
	2nd Trimester	0.37 - 3.60			
	3rd Trimester	0.38 - 4.04			

( References range recommended by the American Thyroid Association) Comments:

- 1. During pregnancy, Free thyroid profile (FT3, FT4 & TSH) is recommended.
- 2. TSH levels are subject to circadian variation, reaches peak levels between 2-4 AM and at a minimum between 6-10 PM. The variation of the day has influence on the measured serum TSH concentrations.

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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

	LIVER FUNCTION TEST(LFT)						
Sample Type : SERUM							
TOTAL BILIRUBIN	0.64	mg/dl	0.3 - 1.2	JENDRASSIK & GROFF			
CONJUGATED BILIRUBIN	0.10	mg/dl	0 - 0.2	DPD			
UNCONJUGATED BILIRUBIN	0.54	mg/dl		Calculated			
S.G.O.T	22	U/L	< 35	KINETIC WITHOUT P5P- IFCC			
S.G.P.T	17	U/L	< 35	KINETIC WITHOUT P5P- IFCC			
ALKALINE PHOSPHATASE	89	U/L	30 - 120	IFCC-AMP BUFFER			
TOTAL PROTEINS	7.6	gm/dl	6.6 - 8.3	Biuret			
ALBUMIN	3.8	gm/dl	3.5 - 5.2	BCG			
GLOBULIN	3.8	gm/dl	2.0 - 3.5	Calculated			
A/G RATIO	1.00			Calculated			



Approved By :

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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

		LIPID I	'RO	FILE				
Sample Type : SERUM	]							
TOTAL CHOLESTER	DL	165		mg/dl		Refere Table B	elow	Cholesterol oxidase/peroxidase
H D L CHOLESTEROI		36		mg/dl		>40		Enzymatic/ Immunoinhibiton
L D L CHOLESTEROL		91.2		mg/dl		Refere Table B	elow	Enzymatic Selective Protein
TRIGLYCERIDES		189		mg/dl		See Table		GPO
VLDL		37.8		mg/dl	mg/dl 15 - 30			Calculated
T. CHOLESTEROL/ HI	DL RATIO	4.58			Refere Table Below		elow	Calculated
TRIGLYCEIDES/ HDL	RATIO	5.25		Ratio		< 2.0		Calculated
NON HDL CHOLESTE	EROL	129		mg/dl		< 130		Calculated
Interpretation								
NATIONAL LIPID ASSO RECOMMENDATIONS (N		TOTAL CHOLESTER	20L	TRIGLYCE	RIDE	LDL CHOLESTEROL	NON HE CHOLESTE	DL ROL
Optimal		<200		<150		<100	<130	
Above Optimal		-		-		100-129	130 - 15	
Borderline High		200-239	_	150-19		130-159	160 - 18	
High Very High		>=240		200-49 >=500		160-189 >=190	190 - 21 >=220	
	Cholesterol : HD	- I Ratio		>=300	,	2-170	>=220	,
Low risk	3.3-4.4							
Average risk	4.5-7.1							
Moderate risk	7.2-11.0							

#### Note:

High risk

>11.0

1. 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

2. NLA-2014 identifies Non HDL Cholesterol (an indicator of all atherogenic lipoproteins such as LDL , VLDL, IDL, Lpa, 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

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Dr. Sumalatha MBBS,DCP Consultant Pathologist





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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method		

HBA1C					
Sample Type : WHOLE BLOOD EDTA					
HBA1c RESULT	5.7	%	Normal Glucose tolerance (non-diabetic): <5.7% Pre-diabetic: 5.7-6.4% Diabetic Mellitus: >6.5%	HPLC	
ESTIMATED AVG. GLUCOSE	117	mg/dl			

Note:

1. Since HbA1c reflects long term fluctuations in the blood glucose concentration, a diabetic patient who is recently under good control may still have a high concentration of HbA1c. Converse is true for a diabetic previously under good control but now poorly controlled .

2. Target goals of < 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targeting a goal of < 7.0 % may not be appropriate.

HbA1c provides an index of average blood glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control .

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Test NameResultUnitBiological Ref. RangeMethod					

<b>BLOOD UREA NITROGEN (BUN)</b>					
Sample Type : Serum					
SERUM UREA	13	mg/dL	13 - 43	Urease GLDH	
Blood Urea Nitrogen (BUN)	6.1	mg/dl	5 - 25	GLDH-UV	

### Increased In:

Impaired kidney function, Reduced renal blood flow {CHF, Salt and water depletion, (vomiting, diarrhea, diuresis, sweating), Shock}, Any obstruction of urinary tract, Increased protein catabolism, AMI, Stress

#### Decreased In:

Diuresis (e.g. with over hydration), Severe liver damage, Late pregnancy, Infancy, Malnutrition, Diet (e.g., low-protein and high-carbohydrate, IV feedings only), Inherited hyperammonemias (urea is virtually absent in blood)

#### Limitations:

Urea levels increase with age and protein content of the diet.

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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Test NameResultUnitBiological Ref. RangeMethod					

	FBS (GLUC	OSE FASTING)					
Sample Type : FLOURIDE PLASMA							
FASTING PLASMA GLUCOSE	100	mg/dl	70 - 100	HEXOKINASE			
INTERPRETATION:							
Increased In							
Diabetes Mellitus							
<ul> <li>Stress (e.g., emotion, burns, shock</li> </ul>	anesthesia)						
<ul> <li>Acute pancreatitis</li> </ul>							
Chronic pancreatitis							
<ul> <li>Wernicke encephalopathy (vitamin E</li> </ul>	31 deficiency)						
• Effect of drugs (e.g. corticosteroids	estrogens, alcoho	I, phenytoin, thiazi	des)				
Decreased In							
Pancreatic disorders							
<ul> <li>Extrapancreatic tumors</li> </ul>							
Endocrine disorders							
Malnutrition							
<ul> <li>Hypothalamic lesions</li> </ul>							
Alcoholism							
<ul> <li>Endocrine disorders</li> </ul>							



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DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method	

PPBS (POST PRANDIAL GLUCOSE)							
Sample Type : FLOURIDE PLASMA							
POST PRANDIAL PLASMA GLUCOSE	123	mg/dl	<140	HEXOKINASE			
INTERPRETATION:							
Increased In  Diabetes Mellitus  Stress (e.g., emotion, burns, shock, anesthe Acute pancreatitis Chronic pancreatitis Wernicke encephalopathy (vitamin B1 deficie Effect of drugs (e.g. corticosteroids, estrogen Decreased In Pancreatic disorders Extrapancreatic tumors Endocrine disorders Malnutrition Hypothalamic lesions	ency)	ytoin, thiazides)					
<ul><li>Alcoholism</li><li>Endocrine disorders</li></ul>							



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DEPARTMENT OF BIOCHEMISTRY							
Test Name	Test NameResultUnitBiological Ref. RangeMethod						

	SERUM C	REATININE		
Sample Type : SERUM				
SERUM CREATININE	0.55	mg/dl	0.51 - 0.95	KINETIC-JAFFE
Increased In:				

- Diet: ingestion of creatinine (roast meat), Muscle disease: gigantism, acromegaly,
- Impaired kidney function.

### Decreased In:

- Pregnancy: Normal value is 0.4-0.6 mg/dL. A value >0.8 mg/dL is abnormal and should alert the clinician to further diagnostic evaluation.
- Creatinine secretion is inhibited by certain drugs (e.g., cimetidine, trimethoprim).

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DE	PARTMENT O	F BIOCHEMI	ISTRY	
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method

	GGT (GA	MMA GLUTA	MYL TRANSP	PEPTIDASE)	
Sample Type : SERUM					
GGT		28	U/L	0 - 55.0	KINETIC-IFCC
INTERPRETATION:					

GGT functions in the body as a transport molecule, helping to move other molecules around the body. It plays a significant role in helping the liver metabolize drugs and other toxins. Increased GGT include overuse of alcohol, chronic viral hepatitis, lack of blood flow to the liver, liver tumor, cirrhosis, or scarred liver, overuse of certain drugs or other toxins, heart failure, diabetes, pancreatitis, fatty liver disease.

Verified By : GOPI



Approved By :

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Dr. Sumalatha MBBS,DCP Consultant Pathologist

CONTACT US



Visit ID	: YGT41532	UHID/MR No	: YGT.0000041384
Patient Name	: Mrs. BANDELA RAJYA LAKSHMI	Client Code	: 1409
Age/Gender	: 27 Y 4 M 14 D /F	Barcode No	: 10816264
DOB	: 12/Jul/1996	Registration	: 25/Nov/2023 09:14AM
Ref Doctor	: SELF	Collected	: 25/Nov/2023 09:21AM
Client Name	: MEDI WHEELS	Received	: 25/Nov/2023 09:42AM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 25/Nov/2023 10:48AM
Hospital Name	:		

DEI	PARTMENT O	F BIOCHEMI	ISTRY	
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method

	UF	RIC ACID -SERUM	[	
Sample Type : SERUM				
SERUM URIC ACID	4	2 mg/dl	2.6 - 6.0	URICASE - PAP

Uric acid is the final product of purine metabolism in the human organism. Uric acid measurements are used in the diagnosis and treatment of numerous renal and metabolic disorders, including renal failure, gout, leukemia, psoriasis, starvation or other wasting conditions, and of patients receiving cytotoxic drugs.

Verified By :



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DEI	PARTMENT O	F BIOCHEMI	STRY	
Test Name	Result	Unit	<b>Biological Ref. Range</b>	Method

	<b>BUN/CREAT</b>	<b>ININE RATIO</b>		
Sample Type : SERUM				
Blood Urea Nitrogen (BUN)	6.1	mg/dl	5 - 25	GLDH-UV
SERUM CREATININE	0.55	mg/dl	0.51 - 0.95	KINETIC-JAFFE
BUN/CREATININE RATIO	11.00	Ratio	6 - 25	Calculated



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	2D ECHO DOPPLER STUDY
MITRAL VALVE	: Normal
AORTIC VALVE	: Normal
TRICUSPID VALVE	: Normal
PULMONARY VALVE	: Normal
RIGHT ATRIUM	: Normal
RIGHT VENTRICLE	: Normal
LEFT ATRIUM	: 2.3 cms
LEFT VENTRICLE	: EDD : 4.3 cm IVS(d) : 0.9cm LVEF : 74 % ESD : 2.4 cm PW (d) : 0.9cm FS : 43 No RWMA
IAS	: Intact
IVS	: Intact
AORTA	: 2.2 cms
PULMONARY ARTERY	: Normal
PERICARDIUM	: Normal
IVS/ SVC/ CS	: Normal
PULMONARY VEINS	: Normal
INTRA CARDIAC MASSE	S: No

Verified By :



Approved By :

Dr.B.Nagaraju MD(Internal Medicine) DN(CARDIOLOGY) APNC Reg.No 70760



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DOPPLER STUDY :				
MITRAL FLOW	: E - 1.7m/sec, A - 0.5m/sec.			
AORTIC FLOW	: 1.2m/sec			
PULMONARY FLOW	: 1.1m/sec			
TRICUSPID FLOW	: TRJV : 1.8 m/sec, RVSP -28 mmHg			
COLOUR FLOW MAPPING:				
IMPRESSION :				
<ul> <li>* NORMAL SIZED CARDIAC CHAMBERS</li> <li>* NO RWMA OF LV</li> <li>* GOOD LV FUNCTION</li> <li>* NORMAL LV FILLING PATTERN</li> <li>* NO MR/NO AR/NO PR</li> <li>* NO TR/NO PAH</li> <li>* NO PE / CLOT /VEGETATIONS.</li> </ul>				

Verified By : GOPI

Approved By :

Dr.B.Nagaraju MD(Internal Medicine) DN(CARDIOLOGY) APNC Reg.No 70760



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Result

<b>DEPARTMENT OF</b>	F CLINICAL PATHOLOGY
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**Test Name** 

Unit

**Biological Ref. Range** 

Method

CUE (COMPLETE URINE EXAMINATION)				
Sample Type : SPOT URINE				
PHYSICAL EXAMINATION				
TOTAL VOLUME	25 ML	ml		
COLOUR	PALE YELLOW			
APPEARANCE	CLEAR			
SPECIFIC GRAVITY	1.010		1.003 - 1.035	Bromothymol Blue
CHEMICAL EXAMINATION				
pH	6.0		4.6 - 8.0	Double Indicator
PROTEIN	NEGATIVE		NEGATIVE	Protein - error of Indicators
GLUCOSE(U)	NEGATIVE		NEGATIVE	Glucose Oxidase
UROBILINOGEN	NEGATIVE	mg/dl	< 1.0	Ehrlichs Reaction
KETONE BODIES	NEGATIVE		NEGATIVE	Nitroprasside
BILIRUBIN - TOTAL	NEGATIVE		Negative	Azocoupling Reaction
BLOOD	NEGATIVE		NEGATIVE	Tetramethylbenzidine
LEUCOCYTE	NEGATIVE		Negative	Azocoupling reaction
NITRITE	NEGATIVE		NEGATIVE	Diazotization Reaction
MICROSCOPIC EXAMINATION	·			·
PUS CELLS	1-2	cells/HPF	0-5	
EPITHELIAL CELLS	3-4	/hpf	0 - 15	
RBCs	NIL	Cells/HPF	Nil	
CRYSTALS	NIL	Nil	Nil	
CASTS	NIL	/HPF	Nil	
BUDDING YEAST	NIL		Nil	
BACTERIA	NIL		Nil	
OTHER	NIL			

Verified By :



Approved By :

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Dr. Sumalatha MBBS,DCP **Consultant Pathologist** 



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Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 25/Nov/2023 01:27PM
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## DEPARTMENT OF CYTOPATHOLOGY

### **PAP SMEAR - CONVENTIONAL**

PAP SMEAR

Lab Ref. No.: YLLD/C- 154/23

Date of Receiving: 25/11/23

SYSTEM: BETHESDA 2014

SPECIMEN: ONE CERVICAL SMEAR. FIXED IN ALCOHOL

ADEQUACY: SATISFACTORY FOR EVALUATION.

**MICROSCOPY**: Smears show predominantly superficial and intermediate squamous epithelial cells seen with inflammatory cells. Epithelial cells show normal nuclear-cytoplasmic ratio. No koilocytosis seen. No evidence of dysplasia/malignancy is seen in the smears examined.

IMPRESSION: NILM (Negative for intraepithelial lesion and malignancy).

### **ASCO/ CAP GUIDELINES :**

	HPV Unknown	HPV Positive	HPV Negative
Unsatisfactory	Repeat cytology after 2- 4 mths	Colposcopy	Repeat cytology after 2- 4 mths
NILM with EC/TZ	Routine screening	HPV genotyping/ repeat co-testing @ 1 Year	Routine Screening
NILM without EC/TZ	HPV teting	Repeat co-testing @ 1 Year	Routine Screening
ASCUS	HPV teting	Colposcopy	Routine Screening
LSIL	Colposcopy	Colposcopy	Repeat cotesting @ 3 year
ASC - H	Colposcopy	Colposcopy	Colopscopy
HSIL	Immediate LEEP	Immediate LEEP	Immediate LEEP
AGC	EB & Endocervical Bx	EB & Endocervical Bx	EB & Endocervical Bx

SCREENING GUIDELINE: 21-29 Years - Cytology only every 3 years ; <21 & 65 yrs - Screening not recommended

Comments- Pap Test is a screening test for cervical cancer. False negativity may be due to inherent limitation of this technique.

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DEPARTMENT OF CYTOPATHOLOGY

## \*\*\* End Of Report \*\*\*

Verified By :



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Dr. Sumalatha MBBS,DCP Consultant Pathologist

