UNITED HOSPITAL
Care Par Excellence

DEPARTMENT OF LABORATORY MEDICINE

Patient Name	: Mrs. SREEDEVI KURNI	Order No	: 1000082628
UHID	: UHJA24000437	Registered On	: 13/04/2024 09:38:30 AM
Age/Sex	: 41/Years Female	Collected On	: 13/04/2024 10:05:09 AM
Ward / Bed No	:	Reported On	: 13/04/2024 02:00:50 PM
Reference	: Dr. Preventive Health Check Up	Bill No	: OPBJA240000573
Station	: At Hospital	Mobile No	: 9480400587
Payer Name	: Mediwheel	Report Status	: Final Report

Test Name	Result	Unit	Bio. Ref. Interval
	BIOCHEMISTRY		
FASTING GLUCOSE (Method: Hexokinase)	95	mg/dL	ADA Guidelines < 100 mg/dl - Normal 100 to 125 mg/dl - Prediabetes ≥ 126 mg/dl - Diabetes
POST PRANDIAL GLUCOSE (Method: Hexokinase)	119	mg/dL	70-140
GLYCOSYLATED HAEMOGLOBIN (HBA1C)			Sample: Whole blood (EDTA)
HBAIC (Method: HPLC)	6.3	%	ADA Guidelines < 5.7% - Normal 5.7 to 6.4% - Prediabetes ≥ 6.5% - Diabetes
Estimated Average Glucose (eAG) (Method: Calculated)	134.11	mg/dL	
THYROID PROFILE (TOTAL T3, TOTAL T4 & TSH)			Sample: Serum
TOTAL T3 (Method:CLIA)	1.13	ng/mL	0.87-1.78
TOTAL T4 (Method:CLIA)	13.98	≈g/dL	5.1-14.1
THYROID STIMULATING HORMONE (TSH) (Method:CLIA: Ultra-sensitive)	1.82	ı IU/mL	0.34 - 5.60 ı IU/mL (Non Pregnant) 0.3 - 4.5 ı IU/mL (I trimester) 0.5 - 5.2 ı IU/mL (II & III trimester)
LIPID PROFILE			Sample: Serum
TOTAL CHOLESTEROL (Method:CHOD-POD)	199	mg/dL	ATP III Guidelines < 200 - Desirable 200-239 - Borderline high ≥ 240 - High
TRIGLYCERIDES (Method:Enzymatic GPO-POD)	82	mg/dL	< 150 - Normal 150-199 - Borderline High 200-499 - High ≥ 500 - Very High
HDL CHOLESTEROL (Method:ENZYMATIC METHOD)	50.4	mg/dL	< 40 - Low ≥ 60 - High

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LDL CHOLESTEROL (Method:ENZYMATIC METHOD)	132.2	mg/dL	<100 - Optimal 100-129 - Near or above optimal 130-159 - Borderline high 160-189 - High ≥190 - Very high
VLDL CHOLESTEROL (Method: Calculated)	16.39	mg/dL	< 30
TOTAL CHOLESTEROL : HDL RATIO (Method: Calculated)	3.9		Low Risk: 3.3 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0
LDL/HDL CHOLESTEROL RATIO (Method: Calculated)	2.6		< 2.5 Optimal
NON HDL CHOLESTEROL (Method: Calculated)	148.6	mg/dL	< 130
URIC ACID (Method:Uricase - POD(Enzymatic))	5.4	mg/dL	2.6-6.0
LIVER FUNCTION TEST			Sample: Serum
TOTAL BILIRUBIN (Method:Dichlorophenyl Diazotization)	0.65	mg/dL	0.3-1.2
DIRECT BILIRUBIN (Method:Dichlorophenyl Diazotization)	0.14	mg/dL	0.0-0.2
INDIRECT BILIRUBIN (Method: Calculated)	0.52	mg/dL	0.2-1.0
TOTAL PROTEIN (Method:BIURET)	7.2	g/dL	6.6-8.3
ALBUMIN (Method:BCG)	3.93	g/dL	3.5-5.2
GLOBULIN (Method: Calculated)	3.27	g/dL	2.3-3.5
AG RATIO (Method: Calculated)	1.20		2:1
SERUM SGOT (Method:IFCC without P5P)	21	U/L	< 35

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SERUM SGPT (Method:IFCC without P5P)	28	U/L	< 35
ALKALINE PHOSPHATASE, SERUM (Method:PNPP AMP Buffer)	86	U/L	46-122
GGT (Method:IFCC)	20	U/L	< 38
UREA (Method:Urease GLDH - Kinetic)	19.3	mg/dL	17-43
BUN/CREATININE RATIO			Sample: Serum
BLOOD UREA NITROGEN(BUN) (Method:Urease GLDH - Kinetic)	9	mg/dL	7.93-20.07
CREATININE (Method:Modified Jaffe, Kinetic)	0.6	mg/dL	0.6-1.1
BUN/CRE-RATIO (Method: Calculated)	15.2		12¯20 : 1

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HAEMATOLOGY

COMPLETE BLOOD COUNT(CBC) — Sample: Whole blood (EDTA)

Test Name	Result	Unit	Bio. Ref. Interval
HAEMOGLOBIN (Method:Photometric Measurement: Oxyhemoglobin method)	12.02	g/dL	12-16
PACKED CELL VOLUME/HEMATOCRIT (PCV/HCT) (Method: Calculated)	37.5	%	37-47
TOTAL WBC COUNT (TLC) (Method:Coulter Principle)	8430	Cells/Cum	4000-11000
DIFFERENTIAL COUNT			
NEUTROPHILS (Method:Optical/Impedance)	65.16	%	40-75
LYMPHOCYTES (Method:Optical/Impedance)	27.08	%	20-45
EOSINOPHILS (Method:Optical/Impedance)	2.29	%	0-6
MONOCYTES (Method:Optical/Impedance)	5.24	%	2-10
BASOPHILS (Method:Optical/Impedance)	0.23	%	0-2
RED BLOOD CORPUSCLES(RBC) (Method:Coulter Principle)	5.12	million/cum	4.0-5.2
MCV (Method:Derived from RBC Histogram)	73.3	fL	78-100
MCH (Method: Calculated)	23.5	pg	27-31
MCHC (Method: Calculated)	32.1	g/dL	31-37
RDW - CV (Method: Calculated)	17.4	%	11.5-14.5
PLATELET COUNT (Method:Electrical Impedance)	2.06	Lakhs/Cum	1.5-4.5

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Test Name	Result	Unit	Bio. Ref. Interval
MEAN PLATELET VOLUME(MPV) (Method:Derived from PLT Histogram)	11.01	fl	9-13
PLATELET DISTRIBUTION WIDTH (PDW) (Method: Calculated)	28.9	fl	9-19
ERYTHROCYTE SEDIMENTATION RATE(ESR) (Method:Modified Westergren Method)	12	mm/hour	1-20

BLOOD GROUPING & RH TYPING — Sample: Whole blood (EDTA)

ABO Group (Method:Agglutination Method)	B		
Rh Factor (Method:Agglutination Method)	Positive		

Interpretation Notes

Note: Both forward and reverse grouping performed

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CLINICAL PATHOLOGY

Sample: Urine

Test Name	Result	Unit	Bio. Ref. Interval
URINE EXAMINATION, ROUTINE			
PHYSICAL EXAMINATION			
VOLUME	25	mL	
COLOUR	Pale Yellow		
APPEARANCE	Clear		
PH	6.0		5.0-8.0
SPECIFIC GRAVITY	1.025		1.005-1.030
CHEMICAL EXAMINATION			
PROTEIN (Method:Protein Error of pH Indicator)	Absent		Absent
GLUCOSE (Method:GOD-POD)	Absent		Absent
KETONE BODIES (Method:Nitroprusside method/ Rothera's test)	Absent		Absent
BILIRUBIN (Method:DIAZO/FOUCHET'S TEST)	Negative		Negative
BILE SALT (Method:Hay's sulfur test)	Absent		Absent
NITRITE (Method:Griess method)	Negative		Negative
UROBILINOGEN (Method:Azo coupling method)	Normal		
LEUKOCYTE ESTERASE (Method:Leukocyte Esterase activity)	Negative		Negative
BLOOD (Method:Peroxidase Reaction)	Negative		Negative
MICROSCOPIC EXAMINATION			

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Test Name	Result	Unit	Bio. Ref. Interval
EPITHELIAL CELLS	2-4	/HPF	0-5
PUS CELLS	0-2	/HPF	0-5
RBCs	Nil	/HPF	0-2
CASTS	Nil	/LPF	
CRYSTALS	Nil		
OTHERS	Nil		
URINE SUGAR, FASTING (Method:GOD-POD)	Absent		
URINE SUGAR (POST PRANDIAL)	Absent		

Verified By
PRAVEEN T

---End of Report---

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*NABL renewal under process.

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DEPARTMENT OF LABORATORY MEDICINE

Patient Name	: Mrs. SREEDEVI KURNI	Order No	: 1000082631
UHID	: UHJA24000437 \	Registered On	: 13/04/2024 09:38:29 AM
Age/Sex	: 41/Years Female	Collected On	: 13/04/2024 04:19:07 PM
Ward / Bed No	:	Reported On	: 15/04/2024 04:44:07 PM
Reference	: Dr. Preventive Health Check Up	Bill No	: OPBJA240000573
Station	: At Hospital	Mobile No	: 9480400587
Payer Name	: Mediwheel	Report Status	: Final Report

Samples

CERVICAL SMEAR - 13/04/2024 04:19 PM

Test Name :PAP SMEAR

NUMBER OF SLIDES RECEIVED: 02
TYPE OF THE SMEAR: Conventional
SOURCE OF THE SMEAR: Ecto and endocervix
CLINICAL DETAILS: Asymptomatic

SPECIMEN ADEQUACY:
Satisfactory for evaluation.
Transformation zone/ Endocervical cell component is present

MICROSCOPY:
Smears show predominantly superficial, intermediate squamous and endocervical cells.
Background shows neutrophils

IMPRESSION: NEGATIVE FOR INTRAEPITHELIAL LESION OR MALIGNANCY (NILM)

DISCLAIMER
1. PLEASE NOTE PAPANICOLAOU SMEAR STUDY IS A SCREENING PROCEDURE FOR CERVICAL CANCER WITH INHERENT FALSE NEGATIVE RESULTS, HENCE SHOULD BE INTERPRETED WITH CAUTION.
2. NO CYTOLOGIC EVIDENCE OF HPV INFECTION IN THE SMEARS STUDIED.

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