UNITED HOSPITAL
Care Par Excellence
A Unit of United Brotherss Healthcare Services Private Limited
No. 110 (30), Madhavan Park Circle, 10th Main Road, 3rd Block, Jayanagar, Bangalore – 560011. T: 080 4566 6666/ 080 6933 3333 E: info@unitedhospital.in

DEPARTMENT OF RADIODIAGNOSIS

Name	Aherwar Nitin Kumar	**Date**	23/03/24
Age	54 years	**Hospital ID**	UHJA23021016
Sex	Male	**Ref.**	Healthcheck

RADIOGRAPH OF THE CHEST (PA – VIEW)

FINDINGS:

Bilateral lung fields are normal.

Bilateral costo-phrenic angles are normal.

Cardia and mediastinal contours are normal.

The bony thorax is grossly normal.

IMPRESSION:

- **No radiographic abnormality.**

Dr. Elluru Santosh Kumar
Consultant Radiologist

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Disclaimer for Radiology Scans and Procedures :

1) Radiology results should be correlated and interpreted by qualified medical professionals only. In case of any clarification, the referring doctors or patients can contact the reception/respective department/doctor.

2) Radiology results are affected by patient body habitus, food consumption, bowel contents, hydration status, foreign bodies and artifacts.

3) Small renal/ureteric stones, some of the pathologies of bowel, peritoneum and retroperitoneum may not be detected on ultrasound study.

4) Antenatal ultrasound: Maternal body variables, gestational age, fetal position at the time of the scan affects the scanning. Patient should come for review scan if and when recommended. Chromosomal anomalies cannot be diagnosed on ultrasound only. If ultrasound markers indicate high risk for chromosomal anomalies, further evaluation including karyotyping may be needed.

5) Duplicate reports can be provided only upto 30 days from the date of scan/procedure.

6) X-ray is a screening modality and not a diagnostic test. It should be correlated clinically and complemented by other requisite imaging modalities and lab tests. X-ray cannot detect soft tissue injuries (like tendon/ ligament injuries) and small renal/ ureteric stones.

7) All disputes relating to the reports are subject to jurisdiction of courts at Bengaluru city only.

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DEPARTMENT OF LABORATORY MEDICINE

Patient Name	: Mr. AHERWAR NITIN KUMAR	Order No	: 1000079051
UHID	: UHJA23021016	Registered On	: 23/03/2024 09:45:18 AM
Age/Sex	: 54/Years Male	Collected On	: 23/03/2024 10:18:32 AM
Ward / Bed No	:	Reported On	: 23/03/2024 04:19:15 PM
Reference	: Dr. Preventive Health Check Up	Bill No	: OPBJA230026006
Station	: At Hospital	Mobile No	: 9653006160
Payer Name	: Mediwheel	Report Status	: Final Report

Test Name	Result	Unit	Bio. Ref. Interval
	BIOCHEMISTRY		
FASTING GLUCOSE (Method: Hexokinase)	109	mg/dL	ADA Guidelines < 100 mg/dl - Normal 100 to 125 mg/dl - Prediabetes ≥ 126 mg/dl - Diabetes
POST PRANDIAL GLUCOSE (Method: Hexokinase)	120	mg/dL	70-140
GLYCOSYLATED HAEMOGLOBIN (HBA1C)			Sample: Whole blood (EDTA)
HBAIC (Method: HPLC)	5.1	%	ADA Guidelines < 5.7% - Normal 5.7 to 6.4% - Prediabetes ≥ 6.5% - Diabetes
Estimated Average Glucose (eAG) (Method: Calculated)	99.66	mg/dL	
THYROID PROFILE (TOTAL T3, TOTAL T4 & TSH)			Sample: Serum
TOTAL T3 (Method:CLIA)	1.02	ng/mL	0.87-1.78
TOTAL T4 (Method:CLIA)	8.17	≈g/dL	5.1-14.1
THYROID STIMULATING HORMONE (TSH) (Method:CLIA: Ultra-sensitive)	4.66	ı IU/mL	0.34-5.60
LIPID PROFILE			Sample: Serum
TOTAL CHOLESTEROL (Method:CHOD-POD)	247	mg/dL	ATP III Guidelines < 200 - Desirable 200-239 - Borderline high ≥ 240 - High
TRIGLYCERIDES (Method:Enzymatic GPO-POD)	186	mg/dL	< 150 - Normal 150-199 - Borderline High 200-499 - High ≥ 500 - Very High
HDL CHOLESTEROL (Method:ENZYMATIC METHOD)	51.9	mg/dL	< 40 - Low ≥ 60 - High

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Test Name	Result	Unit	Bio. Ref. Interval
LDL CHOLESTEROL (Method:ENZYMATIC METHOD)	157.9	mg/dL	<100 - Optimal 100-129 - Near or above optimal 130-159 - Borderline high 160-189 - High ≥190 - Very high
VLDL CHOLESTEROL (Method: Calculated)	37.20	mg/dL	< 30
TOTAL CHOLESTEROL : HDL RATIO (Method: Calculated)	4.8		Low Risk: 3.3 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0
LDL/HDL CHOLESTEROL RATIO (Method: Calculated)	3.0		< 2.5 Optimal
NON HDL CHOLESTEROL (Method: Calculated)	195.1	mg/dL	< 130
URIC ACID (Method:Uricase - POD(Enzymatic))	6.9	mg/dL	3.5-7.2
BUN/CREATININE RATIO			Sample: Serum
BLOOD UREA NITROGEN(BUN) (Method:Urease GLDH - Kinetic)	10	mg/dL	7.93-20.07
CREATININE (Method:Modified Jaffe, Kinetic)	0.94	mg/dL	0.9-1.3
BUN/CRE-RATIO (Method: Calculated)	12		12¯20 : 1
LIVER FUNCTION TEST			Sample: Serum
TOTAL BILIRUBIN (Method:Dichlorophenyl Diazotization)	0.92	mg/dL	0.3-1.2
DIRECT BILIRUBIN (Method:Dichlorophenyl Diazotization)	0.16	mg/dL	0.0-0.2
INDIRECT BILIRUBIN (Method: Calculated)	0.77	mg/dL	0.2-1.0
TOTAL PROTEIN (Method:BIURET)	6.8	g/dL	6.6-8.3

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Test Name	Result	Unit	Bio. Ref. Interval
ALBUMIN (Method:BCG)	4.56	g/dL	3.5-5.2
GLOBULIN (Method: Calculated)	2.24	g/dL	2.3-3.5
AG RATIO (Method: Calculated)	2.03		2:1
SERUM SGOT (Method:IFCC without P5P)	25	U/L	< 50
SERUM SGPT (Method:IFCC without P5P)	27	U/L	< 50
ALKALINE PHOSPHATASE, SERUM (Method:PNPP AMP Buffer)	79	U/L	50-116
GGT (Method:IFCC)	25	U/L	< 55
PROSTATE SPECIFIC ANTIGEN (PSA) (Method:CLIA)	0.31	ng/mL	< 4.0

Interpretation Notes

Serum PSA concentrations should not be interpreted as absolute evidence for the presence or absence of malignant disease nor should serum PSA be used alone as a screening test for malignant disease. For diagnostic purposes, the results obtained by immunometric assay should always be used in combination with the clinical examinations, patient medical history and other findings. The concentration of PSA in a given specimen, determined with assays from different manufacturers, may not be comparable due to differences in assay methods, calibration, and reagent specificity.

UREA (Method:Urease GLDH - Kinetic)	21.9	mg/dL	17-43

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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)	13.67	g/dL	13.5-17.5
PACKED CELL VOLUME/HEMATOCRIT (PCV/HCT) (Method: Calculated)	41.0	%	42-52
TOTAL WBC COUNT (TLC) (Method:Coulter Principle)	5820	Cells/Cum	4000-11000
DIFFERENTIAL COUNT			
NEUTROPHILS (Method:Optical/Impedance)	69.59	%	40-75
LYMPHOCYTES (Method:Optical/Impedance)	20.26	%	20-45
EOSINOPHILS (Method:Optical/Impedance)	3.60	%	0-6
MONOCYTES (Method:Optical/Impedance)	6.18	%	2-10
BASOPHILS (Method:Optical/Impedance)	0.37	%	0-2
RED BLOOD CORPUSCLES(RBC) (Method:Coulter Principle)	4.40	million/cum	4.5-5.9
MCV (Method:Derived from RBC Histogram)	93.2	fL	78-100
MCH (Method: Calculated)	31.1	pg	27-31
MCHC (Method: Calculated)	33.3	g/dL	31-37
RDW - CV (Method: Calculated)	14.7	%	11.5-14.5

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Test Name	Result	Unit	Bio. Ref. Interval
PLATELET COUNT (Method:Electrical Impedance) Remarks: Results are verified on smear. Kindly correlate clinically.	1.20	Lakhs/Cum	1.5-4.5
MEAN PLATELET VOLUME(MPV) (Method:Derived from PLT Histogram)	11.80	fl	9-13
PLATELET DISTRIBUTION WIDTH (PDW) (Method: Calculated)	16.6	fl	9-19
ERYTHROCYTE SEDIMENTATION RATE(ESR) (Method:Modified Westergren Method)	15	mm/hour	1-20
BLOOD GROUPING & RH TYPING			Sample: Whole blood (EDTA)
ABO Group (Method:Agglutination Gel Method)	B		
Rh Factor (Method:Agglutination Gel Method)	Positive		

Interpretation Notes

Note: Both forward and reverse grouping performed

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CLINICAL PATHOLOGY

URINE EXAMINATION, ROUTINE — Sample: Urine

Test Name	Result	Unit	Bio. Ref. Interval
PHYSICAL EXAMINATION			
VOLUME	25	mL	
COLOUR	Pale Yellow		
APPEARANCE	Clear		
PH	6.5		5.0-8.0
SPECIFIC GRAVITY	1.030		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	0-2	/HPF	0-5
PUS CELLS	0-2	/HPF	0-5
RBCs	Nil	/HPF	0-2
CASTS	Nil	/LPF	
CRYSTALS	Nil		
OTHERS	NA		
URINE SUGAR, FASTING (Method:GOD-POD)	Absent		
URINE SUGAR (POST PRANDIAL)	Absent		

Verified By
NAGARATNA

---End of Report---

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*NABL renewal under process.