

Patient Name : Mrs. KONGE DIVYA

Age/Gender : 34 Y 0 M 0 D /F

DOB :

Ref Doctor : SELF

Client Name : MEDI WHEELS

Client Add : F-701, Lado Sarai, Mehravli, N

Hospital Name

UHID/MR No : YOD.0000578148

Client Code : YOD-DL-0021

Barcode No : 10881681

Received

Registration : 13/Jan/2024 09:44AM

Collected : 13/Jan/2024 09:56AM

Reported : 13/Jan/2024 11:32AM

: 13/Jan/2024 10:20AM

DEPARTMENT OF HAEMATOLOGY				
Test Name Result Unit Biological Ref. Range Method				

ESR (ERYTHROCYTE SEDIMENTATION RATE)					
Sample Type : WHOLE BLOOD EDTA					
ERYTHROCYTE SEDIMENTATION RATE	23	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.

Verified By:



Approved By:

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST



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Test Name Result Unit Biological Ref. Range Method

BLOOD GROUP ABO & RH Typing				
Sample Type : WHOLE BLOOD EDTA				
ABO	A			
Rh Typing	POSITIVE			

Method: Hemagglutination Tube method by forward and reverse grouping

#### COMMENTS:

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

Verified By: Mamatha



Approved By:

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST



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: YOD-DL-0021

: 10881681

CBC(COMPLETE BLOOD COUNT)					
Sample Type : WHOLE BLOOD EDTA					
HAEMOGLOBIN (HB)	12.0	g/dl	12.0 - 15.0	Cyanide-free SLS method	
RBC COUNT(RED BLOOD CELL COUNT)	4.08	million/cmm	3.80 - 4.80	Impedance	
PCV/HAEMATOCRIT	36.0	%	36.0 - 46.0	RBC pulse height detection	
MCV	88.2	fL	83 - 101	Automated/Calculated	
MCH	29.4	pg	27 - 32	Automated/Calculated	
MCHC	33.3	g/dl	31.5 - 34.5	Automated/Calculated	
RDW - CV	12.4	%	11.0-16.0	Automated Calculated	
RDW - SD	40.9	fl	35.0-56.0	Calculated	
MPV	9.2	fL	6.5 - 10.0	Calculated	
PDW	9.6	fL	8.30-25.00	Calculated	
PCT	0.28	%	0.15-0.62	Calculated	
TOTAL LEUCOCYTE COUNT	8,370	cells/ml	4000 - 11000	Flow Cytometry	
DLC (by Flow cytometry/Microscopy)					
NEUTROPHIL	58.3	%	40 - 80	Impedance	
LYMPHOCYTE	31.4	%	20 - 40	Impedance	
EOSINOPHIL	2.9	%	01 - 06	Impedance	
MONOCYTE	6.9	%	02 - 10	Impedance	
BASOPHIL	0.5	%	0 - 1	Impedance	
PLATELET COUNT	3.02	Lakhs/cumm	1.50 - 4.10	Impedance	

Verified By:



Approved By:

DR PRANITHA ANAPINDI MD , CONSULTANT PATHOLOGIST



 Visit ID
 : YOD599152
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 : YOD.0000578148

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Age/Gender : 34 Y 0 M 0 D /F Barcode No : 10881681

DOB : Registration : 13/Jan/2024 09:44AM

Ref Doctor: SELFCollected: 13/Jan/2024 09:56AMClient Name: MEDI WHEELSReceived: 13/Jan/2024 10:51AM

Client Add : F-701, Lado Sarai, Mehravli, N Reported : 13/Jan/2024 12:03PM

Hospital Name :

DEPARTMENT OF BIOCHEMISTRY				
Test Name	Result	Unit	Biological Ref. Range	Method

THYROID PROFILE (T3,T4,TSH)					
Sample Type : SERUM					
T3	1.38	ng/ml	0.60 - 1.78	CLIA	
T4	11.02	ug/dl	4.82-15.65	CLIA	
TSH	3.16	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, malnutrition, 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
- in non-thyroidal illness also.
  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 uIU/ 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.

Verified By:



Approved By:



Page 4 of 16



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: MEDI WHEELS Client Name Received : 13/Jan/2024 10:51AM Client Add : F-701, Lado Sarai, Mehravli, N Reported : 13/Jan/2024 12:03PM

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LIVER FUNCTION TEST(LFT)				
Sample Type : SERUM				
TOTAL BILIRUBIN	0.57	mg/dl	0.3 - 1.2	JENDRASSIK & GROFF
CONJUGATED BILIRUBIN	0.11	mg/dl	0 - 0.2	DPD
UNCONJUGATED BILIRUBIN	0.46	mg/dl		Calculated
AST (S.G.O.T)	20	U/L	< 50	KINETIC WITHOUT P5P- IFCC
ALT (S.G.P.T)	17	U/L	< 50	KINETIC WITHOUT P5P- IFCC
ALKALINE PHOSPHATASE	64	U/L	30 - 120	IFCC-AMP BUFFER
TOTAL PROTEINS	7.2	gm/dl	6.6 - 8.3	Biuret
ALBUMIN	4.2	gm/dl	3.5 - 5.2	BCG
GLOBULIN	3	gm/dl	2.0 - 3.5	Calculated
A/G RATIO	1.40			Calculated

Verified By: Mamatha







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DEPARTMENT OF BIOCHEMISTRY				
Test Name Result Unit Biological Ref. Range Method				

LIPID PROFILE				
Sample Type : SERUM				
TOTAL CHOLESTEROL	167	mg/dl	Refere Table Below	Cholesterol oxidase/peroxidase
H D L CHOLESTEROL	40	mg/dl	> 40	Enzymatic/ Immunoinhibiton
L D L CHOLESTEROL	108.4	mg/dl	Refere Table Below	Enzymatic Selective Protein
TRIGLYCERIDES	93	mg/dl	See Table	GPO
VLDL	18.6	mg/dl	< 35	Calculated
T. CHOLESTEROL/ HDL RATIO	4.18	1	Refere Table Below	Calculated
TRIGLYCEIDES/ HDL RATIO	2.33	Ratio	< 2.0	Calculated
NON HDL CHOLESTEROL	127	mg/dl	< 130	Calculated

Interpretation					
NATIONAL CHOLESTEROL EDUCATION		TOTAL	TRI GLYCERI DE	LDL	NON HDL
PROGRAMME (NCEP)		CHOLESTEROL	THI GET OLITIBL	CHOLESTEROL	CHOLESTEROL
Optimal		<200	<150	<100	<130
Above Optimal		-	-	100-129	130 - 159
Borderline High		200-239	150-199	130-159	160 - 189
High		>=240	200-499	160-189	190 - 219
Very High		-	>=500	>=190	>=220
REMARKS Cholesterol : HDL Ratio					<del>-</del>
Low risk					

Average risk 4.5 - 7.1Moderate risk 7.2-11.0 High risk >11.0

Note:

- 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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DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	Biological Ref. Range	Method	

: YOD.0000578148

HBA1C						
Sample Type : WHOLE BLOOD EDTA						
HBA1c RESULT	5.4	%	Normal Glucose tolerance (non-diabetic): <5.7% Pre-diabetic: 5.7-6.4% Diabetic Mellitus: >6.5%	HPLC		
ESTIMATED AVG. GLUCOSE	108	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.

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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DEPARTMENT OF BIOCHEMISTRY					
Test Name	Result	Unit	Biological Ref. Range	Method	

BLOOD UREA NITROGEN (BUN)						
Sample Type : Serum						
SERUM UREA	14	mg/dL	13 - 43	Urease GLDH		
Blood Urea Nitrogen (BUN)	6.5	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	Result	Unit	Biological Ref. Range	Method	

FBS (GLUCOSE FASTING)					
Sample Type : FLOURIDE PLASMA					
FASTING PLASMA GLUCOSE	84	mg/dl	70 - 100	HEXOKINASE	

#### INTERPRETATION:

#### Increased In

- Diabetes Mellitus
- Stress (e.g., emotion, burns, shock, anesthesia)
- Acute pancreatitis
- Chronic pancreatitis
- Wernicke encephalopathy (vitamin B1 deficiency)
- Effect of drugs (e.g. corticosteroids, estrogens, alcohol, phenytoin, thiazides)

## Decreased In

- Pancreatic disorders
- Extrapancreatic tumors
- Endocrine disorders
- Malnutrition
- Hypothalamic lesions
- Alcoholism
- Endocrine disorders

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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Test Name Result Unit Biological Ref. Range Method					

UHID/MR No

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SERUM CREATININE					
Sample Type : SERUM					
SERUM CREATININE	0.63	mg/dl	0.60 - 1.1	KINETIC-JAFFE	
1		-			

#### 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).

Verified By: Mamatha

SURYADEEP PRATAP



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DOB

Ref Doctor : SELF

Collected : 13/Jan/2024 09:56AM : MEDI WHEELS : 13/Jan/2024 10:51AM Client Name Received : 13/Jan/2024 12:03PM Client Add : F-701, Lado Sarai, Mehravli, N Reported

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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Test Name Result Unit Biological Ref. Range Method					

UHID/MR No

Registration

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: 13/Jan/2024 09:44AM

URIC ACID -SERUM					
Sample Type : SERUM					
SERUM URIC ACID		4.5	mg/dl	2.6 - 6.0	URICASE - PAP

Interpretation

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.

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DEPARTMENT OF BIOCHEMISTRY						
Test Name	Test Name Result Unit Biological Ref. Range Method					

UHID/MR No

: YOD.0000578148

BUN/CREATININE RATIO						
Sample Type : SERUM						
Blood Urea Nitrogen (BUN)	6.5	mg/dl	5 - 25	GLDH-UV		
SERUM CREATININE	0.63	mg/dl	0.60 - 1.1	KINETIC-JAFFE		
BUN/CREATININE RATIO	10.30	Ratio	6 - 25	Calculated		

Mamatha

Verified By:

SURYADEEP PRATAP



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## DEPARTMENT OF RADIOLOGY

# **2D ECHO DOPPLER STUDY**

MITRAL VALVE : Normal

AORTIC VALVE : Normal

TRICUSPID VALVE : Normal

PULMONARY VALVE : Normal

RIGHT ATRIUM : Normal

RIGHT VENTRICLE : Normal

LEFT ATRIUM : 3.0 cms

LEFT VENTRICLE

EDD: 4.3 cm IVS(d):0.9 cm LVEF:66 % ESD: 2.7 cm PW (d):0.9 cm FS :33 %

No RWMA

IAS : Intact

IVS : Intact

AORTA : 2.2cms

PULMONARY ARTERY : Normal

PERICARDIUM : Normal

IVS/ SVC/ CS : Normal

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Hospital Name

## DEPARTMENT OF RADIOLOGY

UHID/MR No

PULMONARY VEINS : Normal

INTRA CARDIAC MASSES: No

DOPPLER STUDY:

: E 0.9 m/sec, A 1.0 m/sec. MITRAL FLOW

**AORTIC FLOW** : 1.2m/sec

**PULMONARY FLOW** : 0.8m/sec

: TRJV :2.3 m/sec, RVSP 27 mmHg TRICUSPID FLOW

COLOUR FLOW MAPPING: TRIVIAL TR

# **IMPRESSION:**

- \* NO RWMA OF LV
- \* NORMAL LV SYSTOLIC FUNCTION
- GRADE I LV DIASTOLIC DYSFUNCTION
- TRIVIAL TR
- \* NO PE / CLOT / PAH

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Received : 13/Jan/2024 01:56PM

Reported : 13/Jan/2024 02:15PM

DEPARTMENT OF CLINICAL PATHOLOGY						
Test Name Result Unit Biological Ref. Range Method						

	CUE (COMPLETE U	RINE EXAMIN	(ATION)	
Sample Type : SPOT URINE				
PHYSICAL EXAMINATION				
TOTAL VOLUME	20 ML	ml		
COLOUR	PALE YELLOW	$\wedge$		
APPEARANCE	HAZY			
SPECIFIC GRAVITY	1.002		1.003 - 1.035	Bromothymol Blue
CHEMICAL EXAMINATION				•
pН	5.5		4.6 - 8.0	Double Indicator
PROTEIN	NEGATIVE		NEGATIVE	Protein - error of Indicators
GLUCOSE(U)	NEGATIVE		NEGATIVE	Glucose Oxidase
UROBILINOGEN	0.1	mg/dl	< 1.0	Ehrlichs Reaction
KETONE BODIES	NEGATIVE	/	NEGATIVE	Nitroprasside
BILIRUBIN - TOTAL	NEGATIVE		Negative	Azocoupling Reaction
BLOOD	POSITIVE		NEGATIVE	Tetramethylbenzidine
LEUCOCYTE	POSITIVE		Negative	Azocoupling reaction
NITRITE	NEGATIVE		NEGATIVE	Diazotization Reaction
MICROSCOPIC EXAMINATION				
PUS CELLS	12-15	cells/HPF	0-5	
EPITHELIAL CELLS	6-7	/hpf	0 - 15	
RBCs	1-2	Cells/HPF	Nil	
CRYSTALS	NIL	Nil	Nil	
CASTS	NIL	/HPF	Nil	
BUDDING YEAST	NIL		Nil	
BACTERIA	NIL		Nil	
OTHER	NIL			

Verified By:

Mamatha

Approved By:

DR PRANITHA ANAPINDI



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DEPARTMENT OF CLINICAL PATHOLOGY					
Test Name Result Unit Biological Ref. Range Met					

Received

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

Verified By:
Mamatha

DR PRANITHA ANAPINDI MD, CONSULTANT PATHOLOGIST



# **EYE GLASS PRESCRIPTION**

Name: My. Bandari Balasai Krighna						
Age :	32	<u> </u>	Empl	oyee ID:	599049	
Gender:	~	1			3/01/24	
(unaided PGP		6/a	6/12			
		SPH	CYL	AXIS	BCVA	
Distance	OD	0.50	-	-	€/6	
	os	0.75			6/6	
Add		N 38 C	6 ons	Sin	ENS TYPE  agle Vision Distance agle Vision Near  ocal bgressive	
Remarks: CV - Normal						
					Strics Pvt./for * Pvt./signapour	

Cour Branches at: KPHB PHASE III I MADINAGUDA I VIZAG

• O40-35353535 ⊕ www.yodadiagnostics.com helpdesk@yodalifeline.in helpdesk@yodalifeline.in helpdesk@yodalifeline.in helpdesk@yodalifeline.in our Branches at: KPHB PHASE III I MADINAGUDA I VIZAG



599049

Mr. Banderi Balasai Krishna 32/M

13/01/2024

Has came for general Eye Escentillo Mo Ho DM and HETN

Melo Pap not bosonglit

Slit lamp ence

5. Elp Wal a Mormal

-- 0/s Los/ 2 Normal

i. cu





DEPARTMENT OF RADIOLOGY							
Patient Name	Mr. BANDARI BALASAI KRISHNA	Visit ID	YOD599049	Registration Date	13-01-2024 08:13 AM		
Age / Gender	32/MALE	UHID	YOD.0000578057	Collection Date	01-01-0001 12:00 AM		
Ref Doctor	SELF	Hospital Name		Received Date			
Barcode	10881544	Sample Type		Reported Date	13-01-2024 09:38 AM		

## **ULTRASOUND WHOLE ABDOMEN**

**Clinical Details:** General check-up.

**LIVER:** Normal in size (123mm) and echo-texture. No focal lesion is seen. Intra hepatic biliary channels are not dilated. Visualized common bile duct & portal vein appears normal.

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

PANCREAS: Normal in size and echotexture. No ductal dilatation. No calcifications / calculi.

SPLEEN: Normal in size (86mm) and echotexture. No focal lesion is seen.

RIGHT KIDNEY: measures 82x34mm. Normal in size and echotexture. Cortico-medullary differentiation well

maintained. No focal lesion is seen. Collecting system does not show any dilatation or calculus.

LEFT KIDNEY: measures 95x48mm. Normal in size and echotexture. Cortico-medullary differentiation well

maintained. No focal lesion is seen. Collecting system does not show any dilatation or calculus.

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

**PROSTATE:** Normal in size and echo-texture, volume: 14cc.

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

No free fluid is seen in peritoneal cavity.

Anterior abdominal wall defects measuring 18x13mm & 10x9mm noted just towards right of umbilicus and supraumbilical regions respectively. Omentum herniating through the defects.

## **IMPRESSION:**

- · Paraumbilical hernias.
- No other significant sonological abnormality detected.

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

Suggested clinical correlation & follow up



Approved by

S. SHRAVAN KUMAR (DNB) CONSULTANT RADIOLOGIST



Yoda Diagnostics Pvt Ltd,



DEPARTMENT OF RADIOLOGY							
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Barcode	10881544	Sample Type		Reported Date	13-01-2024 09:25 AM		

## X-RAY CHEST PA VIEW

# **FINDINGS**:

Trachea is midline.

Mediastinal outline, and cardiac silhouette are normal.

Bilateral lung fields show normal vascular pattern with no focal lesion.

Bilateral hila are normal in density.

Bilateral costo-phrenic angles and domes of diaphragms are normal.

The rib cage and visualized bones appear normal.

# **IMPRESSION:**

· No significant abnormality detected.

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

Suggested clinical correlation & follow up



Approved by

Dr. ANNAREDDY SIVAKALA MBBS, DNB , CONSULTANT RADIOLOGIST



Yoda Diagnostics Pvt Ltd,