

EYE GLASS PRESCRIPTION

Name : Mr. Kongala Sudhir Johnson
 Age : 55 Employee ID: 610542
 Gender: M Date: 27/01/24

Vn ✓
 (unaided)
 PGP

6/60	6/60
------	------

Distance		SPH	CYL	AXIS	BCVA
		OD	+2.50	—	
OS	+2.50	—		6/6p	

Add

+2.50	+2.50
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N6 @ 38 cms

- LENS TYPE**
- Single Vision Distance
 - Single Vision Near
 - Bifocal
 - Progressive
 - UV-Coating

Remarks: CV - Normal



MR. Kongala Sudhix Johnson

27/01/24

55/M

G10542

Has come for general eye exam

H/O DM and HTN since 10 years old

H/O using PGP since 1 year old

slit lamp exam

R.O/D L.O/D L Normal (Lens → NSI)

R.O/S L.O/S L Normal (Lens → NSI)

R.CM L.CM L Normal



610542
55 Years

SUDHIR JOHNSON
Male

27-Jan-24 11:00:33 AM
YODA LIFELINE DIAGNOSTICS

Rate 67 . Sinus Rhythm.....Normal P axis, V-rate 50- 99

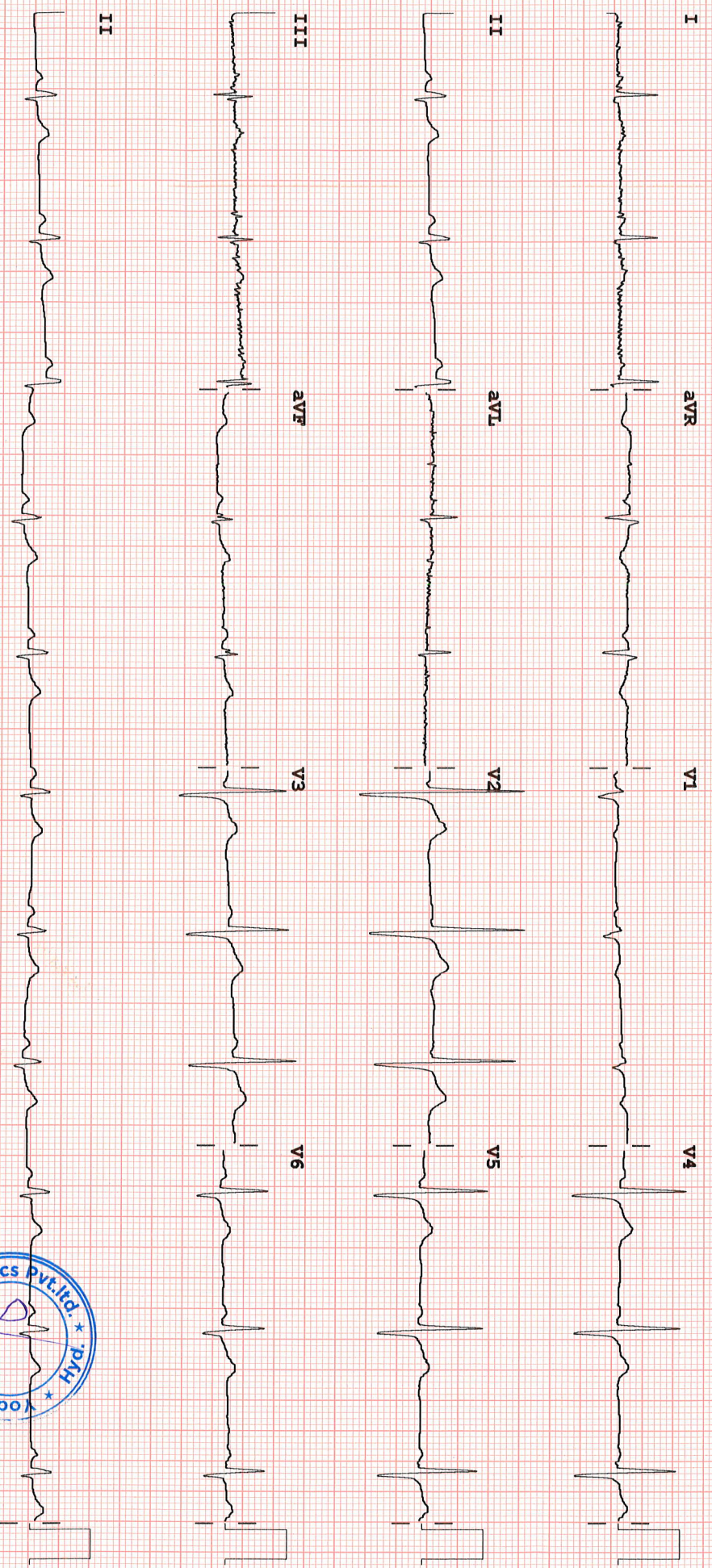
PR 137
QRSD 97
QT 363
QTc 383

--AXIS--
P 51
QRS 28
T 56

12 Lead, Standard Placement

- NORMAL ECG -

Unconfirmed Diagnosis



Device:

Speed: 25 mm/sec

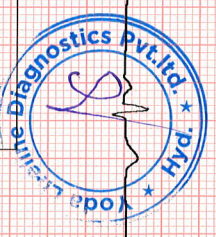
Limb: 10 mm/mV

Chest: 10.0 mm/mV

F 50~ 0.15-100 Hz

100B CL

P?



DEPARTMENT OF RADIOLOGY

Patient Name	Mr. KONGALA SUDHIR JOHNSON	Visit ID	YOD610542	Registration Date	27-01-2024 08:14 AM
Age / Gender	55/MALE	UHID	YOD.0000589096	Collection Date	27-01-2024 08:23 AM
Ref Doctor	SELF	Hospital Name		Received Date	27-01-2024 08:57 AM
Barcode	10899769	Sample Type		Reported Date	27-01-2024 09:14 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

DEPARTMENT OF RADIOLOGY

Patient Name	Mr. KONGALA SUDHIR JOHNSON	Visit ID	YOD610542	Registration Date	27-01-2024 08:14 AM
Age / Gender	55/MALE	UHID	YOD.0000589096	Collection Date	27-01-2024 08:23 AM
Ref Doctor	SELF	Hospital Name		Received Date	27-01-2024 08:57 AM
Barcode	10899769	Sample Type		Reported Date	27-01-2024 10:01 AM

ULTRASOUND WHOLE ABDOMEN

Clinical Details : General check-up.

LIVER: Normal in size (127mm) 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 : Partially 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 97x51mm. 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 103x48mm. 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 : 17.3cc.

No enlarged nodes are visualized. No retro-peritoneal lesion is identified. Great vessels appear normal. No free fluid is seen in peritoneal cavity.

Umbilical hernia noted defect size measuring 1.1cm with omentum as content. Contents are reducible.

IMPRESSION:

- Umbilical hernia
- No other significant sonological abnormality detected.

*** End Of Report ***

Suggested clinical correlation & follow up



Approved by

Dr. ANNAREDDY SIVAKALA
MBBS, DNB , CONSULTANT
RADIOLOGIST



Yoda Diagnostics Pvt Ltd,

Door No: 6-3-862/A, Lal Bungalow add on, Ameerpet, Hyderabad - 500016 helpdesk@yodalifeline.in 040-35353535

Visit ID	: YOD610542	UHID/MR No	: YOD.0000589096
Patient Name	: Mr. KONGALA SUDHIR JOHNSON	Client Code	: YOD-DL-0021
Age/Gender	: 55 Y 0 M 0 D /M	Barcode No	: 10899769
DOB	:	Registration	: 27/Jan/2024 08:14AM
Ref Doctor	: SELF	Collected	: 27/Jan/2024 08:23AM
Client Name	: MEDI WHEELS	Received	: 27/Jan/2024 09:02AM
Client Add	: F-701, Lado Sarai, Mehrauli, N	Reported	: 27/Jan/2024 11:22AM
Hospital Name	:		

DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Unit	Biological Ref. Range	Method
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ESR (ERYTHROCYTE SEDIMENTATION RATE)

Sample Type : WHOLE BLOOD EDTA

ERYTHROCYTE SEDIMENTATION RATE	5	mm/1st hr	0 - 15	Capillary Photometry
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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: Sick cell anemia, spherocytosis, hypofibrinogenemia, or polycythemia vera.

Verified By :
M Thirumalesh Reddy



Approved By :


DR. ABDUL ALEEM MOHAMMED
 MD, DNB (PATHOLOGY)
 Fellowship in Cytogenetics (USA)

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Client Name	: MEDI WHEELS	Received	: 27/Jan/2024 09:02AM
Client Add	: F-701, Lado Sarai, Mehrauli, N	Reported	: 27/Jan/2024 11:26AM
Hospital Name	:		

DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Unit	Biological Ref. Range	Method
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BLOOD GROUP ABO & RH Typing

Sample Type : WHOLE BLOOD EDTA

ABO	AB			
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. Advsiied cross matching before transfusion

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DEPARTMENT OF HAEMATOLOGY


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CBC (COMPLETE BLOOD COUNT)
Sample Type : WHOLE BLOOD EDTA

HAEMOGLOBIN (HB)	13.7	g/dl	13.0 - 17.0	Cyanide-free SLS method
RBC COUNT (RED BLOOD CELL COUNT)	4.79	million/cmm	4.50 - 5.50	Impedance
PCV/HAEMATOCRIT	41.4	%	40.0 - 50.0	RBC pulse height detection
MCV	86.4	fL	83 - 101	Automated/Calculated
MCH	28.6	pg	27 - 32	Automated/Calculated
MCHC	33.1	g/dl	31.5 - 34.5	Automated/Calculated
RDW - CV	12.7	%	11.0-16.0	Automated Calculated
RDW - SD	41	fl	35.0-56.0	Calculated
MPV	9.9	fL	6.5 - 10.0	Calculated
PDW	10.7	fL	8.30-25.00	Calculated
PCT	0.24	%	0.15-0.62	Calculated
TOTAL LEUCOCYTE COUNT	6,080	cells/ml	4000 - 11000	Flow Cytometry
DLC (by Flow cytometry/Microscopy)				
NEUTROPHIL	60.7	%	40 - 80	Impedance
LYMPHOCYTE	27.3	%	20 - 40	Impedance
EOSINOPHIL	2.5	%	01 - 06	Impedance
MONOCYTE	9.2	%	02 - 10	Impedance
BASOPHIL	0.3	%	0 - 1	Impedance
PLATELET COUNT	2.46	Lakhs/cumm	1.50 - 4.10	Impedance

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Client Name : MEDI WHEELS	Received : 27/Jan/2024 08:57AM
Client Add : F-701, Lado Sarai, Mehrauli, N	Reported : 27/Jan/2024 12:20PM
Hospital Name :	

DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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THYROID PROFILE (T3,T4,TSH)

Sample Type : SERUM

T3	1.01	ng/ml	0.60 - 1.78	CLIA
T4	8.77	ug/dl	4.82-15.65	CLIA
TSH	3.25	uIU/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 propranolol 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

(Reference range recommended by the American Thyroid Association)

Comments:

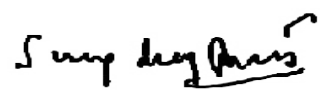
1. During pregnancy, Free thyroid profile (FT3, FT4 & TSH) is recommended.
2. TSH levels are subject to circadian variation, reaches peak levels between 2-4 AM and at a minimum between 6-10 PM. The variation of the day has influence on the measured serum TSH concentrations.

Verified By :

M Thirumalesh Reddy



Approved By :



SURYADEEP PRATAP
Senior Biochemist

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Client Name : MEDI WHEELS	Received : 27/Jan/2024 08:57AM
Client Add : F-701, Lado Sarai, Mehravli, N	Reported : 27/Jan/2024 09:59AM
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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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LIVER FUNCTION TEST(LFT)

Sample Type : SERUM

TOTAL BILIRUBIN	0.53	mg/dl	0.3 - 1.2	JENDRASSIK & GROFF
CONJUGATED BILIRUBIN	0.10	mg/dl	0 - 0.2	DPD
UNCONJUGATED BILIRUBIN	0.43	mg/dl		Calculated
AST (S.G.O.T)	21	U/L	< 50	KINETIC WITHOUT P5P-IFCC
ALT (S.G.P.T)	28	U/L	< 50	KINETIC WITHOUT P5P-IFCC
ALKALINE PHOSPHATASE	65	U/L	30 - 120	IFCC-AMP BUFFER
TOTAL PROTEINS	7.3	gm/dl	6.6 - 8.3	Biuret
ALBUMIN	4.8	gm/dl	3.5 - 5.2	BCG
GLOBULIN	2.5	gm/dl	2.0 - 3.5	Calculated
A/G RATIO	1.92			Calculated

Verified By :
M Thirumalesh Reddy



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SURYADEEP PRATAP
 Senior Biochemist

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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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LIPID PROFILE
Sample Type : SERUM

TOTAL CHOLESTEROL	180	mg/dl	Refere Table Below	Cholesterol oxidase/peroxidase
H D L CHOLESTEROL	45	mg/dl	> 40	Enzymatic/ Immunoinhibiton
L D L CHOLESTEROL	103.4	mg/dl	Refere Table Below	Enzymatic Selective Protein
TRIGLYCERIDES	158	mg/dl	See Table	GPO
VLDL	31.6	mg/dl	< 35	Calculated
T. CHOLESTEROL/ HDL RATIO	4.00		Refere Table Below	Calculated
TRIGLYCEIDES/ HDL RATIO	3.51	Ratio	< 2.0	Calculated
NON HDL CHOLESTEROL	135	mg/dl	< 130	Calculated

Interpretation

NATIONAL CHOLESTEROL EDUCATION PROGRAMME (NCEP)	TOTAL CHOLESTEROL	TRI GLYCERIDE	LDL CHOLESTEROL	NON HDL 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

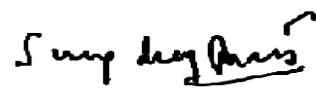
REMARKS	Cholesterol : HDL Ratio
Low risk	3.3-4.4
Average risk	4.5-7.1
Moderate risk	7.2-11.0
High risk	>11.0

Note:

- 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
- 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.
- Apolipoprotein B is an optional, secondary lipid target for treatment once LDL & Non HDL goals have been achieved
- 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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SURYADEEP PRATAP
 Senior Biochemist

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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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PSA (PROSTATE SPECIFIC ANTIGEN) - TOTAL

Sample Type : SERUM

PROSTATE SPECIFIC ANTIGEN	0.43	ng/mL	< 4.0	CLIA
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INTERPRETATION:
 Raised Total PSA levels may indicate prostate cancer, benign prostate hypertation (BPH), or inflammation of the prostate. Prostate manipulation by biopsy or rigorous physical activity may temporarily elevate PSA levels. The blood test should be done before surgery or six weeks after manipulation. The total PSA may be ordered at regular intervals during treatment of men who have been diagnosed with Prostate cancer and in prostatic cancer cases under observation.

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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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HBA1C

Sample Type : WHOLE BLOOD EDTA

HBA1c RESULT	6.4	%	Normal Glucose tolerance (non-diabetic): <5.7% Pre-diabetic: 5.7-6.4% Diabetic Mellitus: >6.5%	HPLC
ESTIMATED AVG. GLUCOSE	137	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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BLOOD UREA NITROGEN (BUN)

Sample Type : Serum

SERUM UREA	19	mg/dL	13 - 43	Urease GLDH
Blood Urea Nitrogen (BUN)	8.9	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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FBS (GLUCOSE FASTING)

Sample Type : FLOURIDE PLASMA

FASTING PLASMA GLUCOSE	125	mg/dl	70 - 100	HEXOKINASE
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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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PPBS (POST PRANDIAL GLUCOSE)

Sample Type : FLOURIDE PLASMA

POST PRANDIAL PLASMA GLUCOSE	151	mg/dl	<140	HEXOKINASE
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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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Ref Doctor	: SELF	Collected	: 27/Jan/2024 08:23AM
Client Name	: MEDI WHEELS	Received	: 27/Jan/2024 08:57AM
Client Add	: F-701, Lado Sarai, Mehravli, N	Reported	: 27/Jan/2024 09:59AM
Hospital Name	:		

DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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SERUM CREATININE

Sample Type : SERUM

SERUM CREATININE	1.16	mg/dl	0.70 - 1.30	KINETIC-JAFFE
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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 :
M Thirumalesh Reddy



Approved By :

SURYADEEP PRATAP
Senior Biochemist

Visit ID	: YOD610542	UHID/MR No	: YOD.0000589096
Patient Name	: Mr. KONGALA SUDHIR JOHNSON	Client Code	: YOD-DL-0021
Age/Gender	: 55 Y 0 M 0 D /M	Barcode No	: 10899769
DOB	:	Registration	: 27/Jan/2024 08:14AM
Ref Doctor	: SELF	Collected	: 27/Jan/2024 08:23AM
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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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GGT (GAMMA GLUTAMYL TRANSPEPTIDASE)

Sample Type : SERUM

GGT	29	U/L	0 - 55.0	KINETIC-IFCC
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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 :
M Thirumalesh Reddy



Approved By :

SURYADEEP PRATAP
Senior Biochemist

Visit ID	: YOD610542	UHID/MR No	: YOD.0000589096
Patient Name	: Mr. KONGALA SUDHIR JOHNSON	Client Code	: YOD-DL-0021
Age/Gender	: 55 Y 0 M 0 D /M	Barcode No	: 10899769
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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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URIC ACID -SERUM

Sample Type : SERUM

SERUM URIC ACID	7.8	mg/dl	3.5 - 7.20	URICASE - PAP
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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.

Verified By :
M Thirumalesh Reddy



Approved By :

SURYADEEP PRATAP
Senior Biochemist

Visit ID	: YOD610542	UHID/MR No	: YOD.0000589096
Patient Name	: Mr. KONGALA SUDHIR JOHNSON	Client Code	: YOD-DL-0021
Age/Gender	: 55 Y 0 M 0 D /M	Barcode No	: 10899769
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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method
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BUN/CREATININE RATIO

Sample Type : SERUM				
Blood Urea Nitrogen (BUN)	8.9	mg/dl	5 - 25	GLDH-UV
SERUM CREATININE	1.16	mg/dl	0.70 - 1.30	KINETIC-JAFFE
BUN/CREATININE RATIO	7.65	Ratio	6 - 25	Calculated

Verified By :
M Thirumalesh Reddy



Approved By :

Suryadeep Pratap
SURYADEEP PRATAP
 Senior Biochemist

Visit ID : YOD610542	UHID/MR No : YOD.0000589096
Patient Name : Mr. KONGALA SUDHIR JOHNSON	Client Code : YOD-DL-0021
Age/Gender : 55 Y 0 M 0 D /M	Barcode No : 10899769
DOB :	Registration : 27/Jan/2024 08:14AM
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Client Name : MEDI WHEELS	Received : 27/Jan/2024 10:16AM
Client Add : F-701, Lado Sarai, Mehravli, N	Reported : 27/Jan/2024 11:21AM
Hospital Name :	

DEPARTMENT OF CLINICAL PATHOLOGY

Test Name	Result	Unit	Biological Ref. Range	Method
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CUE (COMPLETE URINE EXAMINATION)

Sample Type : SPOT URINE

PHYSICAL EXAMINATION

TOTAL VOLUME	20 ML	ml		
COLOUR	PALE YELLOW			
APPEARANCE	CLEAR			
SPECIFIC GRAVITY	1.002		1.003 - 1.035	Bromothymol Blue

CHEMICAL EXAMINATION

pH	5.5		4.6 - 8.0	Double Indicator
PROTEIN	NEGATIVE		NEGATIVE	Protein - error of Indicators
GLUCOSE(U)	POSITIVE (+)		NEGATIVE	Glucose Oxidase
UROBILINOGEN	0.1	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	1-2	/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 :
M Thirumalesh Reddy



Approved By :


DR. ABDUL ALEEM MOHAMMED
 MD, DNB (PATHOLOGY)
 Fellowship in Cytogenetics (USA)

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DEPARTMENT OF CLINICAL PATHOLOGY

Test Name	Result	Unit	Biological Ref. Range	Method
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*** End Of Report ***



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