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# Respected Sir/Madam,

Please find corporate HC appointment details scheduled for 24-06-2023 at your ANANDRISHLII MEDICAL CENTRE LLP - Pune Center.

Points to note:-

Collect photocopy of employee ID proof if health check is through an employer.

Collect photocopy of personal ID proof if health check is for insurance.

Collect MER as per package details & that company's format (alreedy shared).

By 12 noon of appointment date, share Work order number & visit status (Show/No show

Upload reports in Adbhutam portal as per specifications given earlier.

				- Appoint	ment Box	DEPOS CONTRACT		_
Corporate IPW	Agresoment Name	Package nume	Packago Inculsions	Dualikner Marro	ALF	Session Sessione	NEA,	7
ARCOFEMI HEALTHCARE LIMITED	ARCOFEMI MEDIWHEEL MALE AHC CREDIT PAN INDIA OP AGREEMENT	ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS ABOVE 50Y MALE - 2D ECHO - PAN INDIA - FY2324	Doctor URINE GLUCOSE (FASTING), X-RAY-CHEST PALHEMOGRAM + PERIPHERAL SMEARZ D ECHO, PROSTATIC SPECIEIC ANTIGEN (PSA TOTAL), LIVER FUNCTION TEST (LFT), LILTRASOUND-WHOLE ABDOMEN, PERIPHERAL SMEAR, GAMMA GLUTAMYL TRANFERASE (GGT), DIET CONSULTATION, UPLINE GLUCOSE (POST PRANDIÁL), ECG. BCOOD GROUP ABO. AND RH FACTOR, GLUCOSE, POST PRANDIAL), ECG. BCOOD GROUP ABO. AND RH FACTOR, GLUCOSE, POST PRANDIAL (PP), 2 HOURS (POST MEAL), LIPID PROFILE (TOTAL T3, TOTAL T4, TSH), COMPLETE URINE EXAMINATION, GLUCOSE, FASTING. RENAL PROFILE RENAL FUNCTION TEST (RFT/KFT), BODY MASS INDEX (BMI), X-Ray Chest PA. Prostatic Spocific Antigen (PSA Total), Utrasautra - Whole Abdomen, Lipid Profile	MR KUMAR SANJEEV	male	Self	39	

Please login to AHCN Portal for more details.

AHCN Login Url : Click on Link

Regards, Team Clinic Operations Apollo Health and Lifestyle Ltd.,



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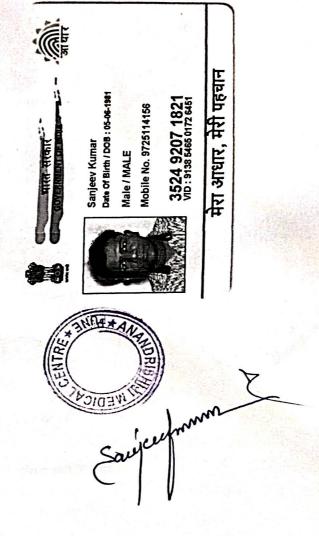




	C ATC
PATIENT MY Same	
AGE YRS	SEX- Hale
HEIGHT- cms	WEIGHT- kg BP- mmhg
166 cm 7	12
OPTHAL CONSULTATION:	RIGHT EYE: - presbyopia LEFT EYE: - Glor vision - O OVERALL EYE CHECKUP: no squint-
PHYSICIAN CONSULTATION:	PAST HISTORY.  no major medical
PRESENT COMPLAINTS-	surfical
no complaints	history
THE CENTRE AND A STATE OF THE S	CNS CNS (A)

UNFIT

Reg. No. 1811070027

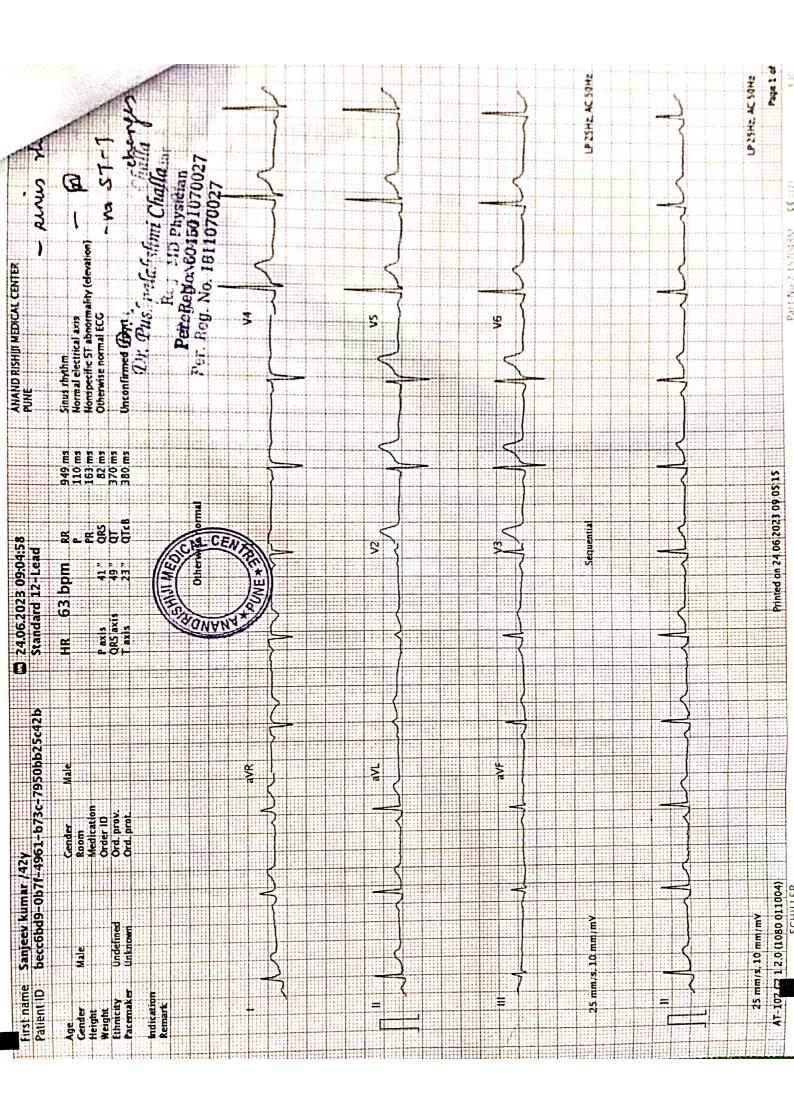






Sanjeev Kumar

Southern





PATIENT'S NAME: MR. SANJEEV KUMAR

REF. CLINICIAN : APOLLO

AGE : 42 Yrs.

DATE: 24-Jun-23

# **Ultrasound Abdomen and Pelvis**

Liver: Normal in position shape and shows diffuse bright Echotexture s/o fatty liver. Span: 13.8 cms.
Hepatic and portal venous radicles are normal.
No IHBR dilatation seen. No focal lesion seen

Spleen: Normal position & echopattern.

Pancreas: Head, body and tail well visualized, bright in echotexture and normal size. No dilatation of main pancreatic duct or focal lesion seen.

Gall bladder: Contracted. (Post prandial Scan) No obvious calculus seen.

Common duct and Portal vein: Normal.

Kidneys:Size:Right kidney: 9.8 x 4.0 cms Left kidney: 11.0 x 5.6 cms Normal position, shape, echo pattern and corticomedullary differentiation seen. No calculi, hydronephrosis or focal parenchymal lesion seen. Moves freely with respiration.

Retroperitoneum: not visualized due to bowel gas.

**Urinary bladder:** Well distended and normal in shape. Wall thickness is within normal limits. No calculus, mural lesion or diverticulum seen.

Prostate: Normal in echopattern and shape. No focal lesion seen.

It measures 3.7 x 3.4 x 3.2 cms (volume- 22 cc). Prostatic calcification seen.

IMPRESSION: Fatty lives

Dr.Trupti S. Jagdale (7.7) Consultant Radiologist Dr. Karuna Agwane Consultant Radiologist

Dr .Mitesh Katariya Consultant Radiologist













PATIENT'S NAME: MR. SANJEEV KUMAR

AGE: 42 Yrs

REF. CLINICIAN : APOLLO

DATE: 24-Jun-23

# 2 DIMENSIONAL ECHOCARDIOGRAPHY & COLOUR DOPPLER REPORT

# M-MODE MEASUREMENTS:

LA	27	mm
AO root	26	mm
LVID(d)	39	mm
LVID (s)	25	mm
IVS (d)	11	mm
LVPW (d)	n many 11 marks and	mm
LVEF	60	%

# **DOPPLER STUDY:**

E wave velocity: 0.51 m/sec A wave velocity: 0.47 m/sec

E/A ratio > 1

	PEAK (mmHg)	GRADE OF REGURGITATI ON
MITRAL	N	Trivial
AORTIC	10	NIL
TRICUSPID	N	Trivial
PULMONA RY	N	Nil



P.T.O



# 2 DIMENSIONAL ECHOCARDIOGRAPHY & COLOUR DOPPLER REPORT

# **COMMENTS:**

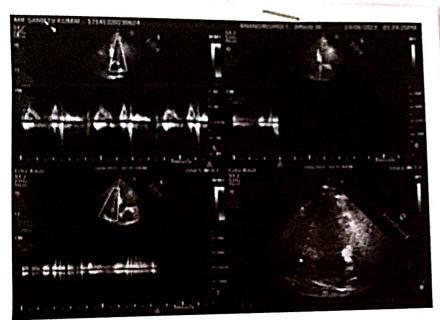
- No LV regional wall motion abnormality at rest.
- Normal resting LV systolic function. LVEF = 60%.
- Normal LV diastolic function.
- Normal chamber dimensions. No LA/LV enlargement.
- Mitral valve normal. Trivial mitral regurgitation.
- Annulo-papillary apparatus appears intact.
- Aortic valve is trileaflet.
- Structurally normal tricuspid valve. Trivial TR.
   PASP by TR jet 25 mmHg. No pulmonary hypertension.
- Normal RV systolic function. IVC normal. IAS & IVS are intact.
- No LV clot/thrombus/pericardial effusion/ vegetation.

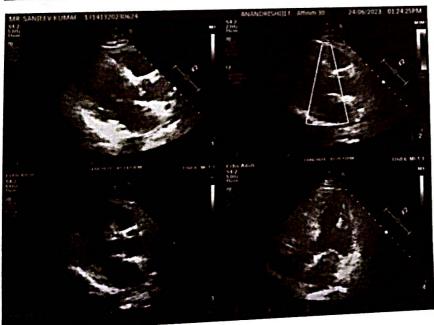
# SUMMARY:

- ➤ Normal LV systolic function. LVEF=60%
- No Regional wall motion abnormality at rest.
- Normal LV diastolic function.
- No pulmonary hypertension. IVC- normal

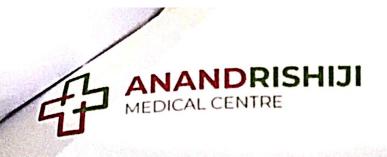
Q1 her

Dr. Shripad Bhivaskar
D.N.B (General Medicine), Dr.N.B. (Cardiology)









Patient's Name	MR. SANJEEV KUMAR	Age/Sex	42Y/ MALE
Ref By	MEDIBUDDY	Date	24/06/2023
ter by			

# X-RAY CHEST PA VIEW

Both the lung fields appear normal.

Both costophrenic angles are normal.

The hila, mediastinal and diaphragmatic outlines appear normal.

The cardiac shadow appears normal.

The bony thoracic cage and soft tissues appear normal.

Impression:- No abnormality detected.

\*Kindly correlate clinically. PLEASE NOTE- Investigations have their limitations. Radiological / Pathological and other investigations never confirm the final Diagnosis. They help in Diagnosing the Disease in correlation to clinical symptom and other related test. Please interpret accordingly and suggest further investigations / Second opinion if clinically indicated. In case of any typing errors please send the report back for correction

DR. RUTUJA DOSHI.

MBBS, DMRE.

Consultant Radiologist.





Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

 Ref. By Dr
 : APOLLO
 Sample Coll. Date
 : 24-Jun-2023
 8:37 AM

 Patient ID
 : 062324012
 Authentication Date
 : 24-Jun-2023
 4:57 PM

 Sample Coll By
 : ANANDRISHIJI MEDICAL CENTRE
 Report Date
 : 24-Jun-2023
 3:59 PM

### **CLINICAL PATHOLOGY**

Investigation	Result	Unit	Bio. Ref. Interval	
URINE EXAMINATION				
PHYSICAL EXAMINATION				
COLOUR	Pale Yellow		Pale Yellow	
APPEARANCE	Clear		Clear	
PH	6.0		5.0-7.5	
SPECIFIC GRAVITY	1.005		1.002-1.030	
CHEMICAL EXAMINATION				
PROTIENS	Absent		Negative	
GLUCOSE	Absent		Negative	
KETONE BODIES	Absent		Negative	
BILLIRUBIN	Absent		Negative	
BLOOD	Absent		Negative	
NITRITE	Absent		Negative	
MICROSCOPIC EXAMINATION				
PUS CELLS	Occasional	/ HPF	0-5	
RED BLOOD CELLS	Absent	/ HPF	Nil	
EPITHELLIAL CELLS	Occasional	/ HPF	< 10	
CASTS	Absent		Absent	
CRYSTALS	Absent		Absent	
YEAST CELLS	Absent		Absent	
BACTERIA	Absent		Absent	
MUCUS THREADS	Absent		Absent	
TRICHOMONAS VAGINAILS	Absent		Absent	
SPERMATOZA	Absent		Absent	
LEUKOCYTES	Absent	ng/ml		
DEPOSIT	Absent		Absent	







Patient Name : MR. SANJEEV KUMAR

Age / Gender : 42 Years / Male

Ref. By Dr : APOLLO
Patient ID : 062324012

Sample Coll By :ANANDRISHIJI MEDICAL CENTRE

Client Name : APOLLO

Registration Date : 24-Jun-2023 8:37 AM

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Authentication Date : 24-Jun-2023 4:57 PM

**Report Date** : 24-Jun-2023 4:55 PM

# **GLUCOSE FASTING, PLASMA**

Investigation	Result	Unit	Bio. Ref. Interval	
BLOOD SUGAR FASTING	96.2	mg/dL	74-106	
METHOD	Hexokinase	9		

# Interpretation:

The fasting (F) blood glucose test is the test most commonly used to diagnose diabetes. It measures blood glucose levels after a period of fasting, usually at least eight hours without food or liquid (except water). This test is more definitive than a random test, because there is no chance that it has been influenced by recent food intake

COMMENT Please correlate with clinical condition







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

Ref. By Dr: APOLLOSample Coll. Date: 24-Jun-20238:37 AMPatient ID: 062324012Authentication Date: 24-Jun-20234:57 PMSample Coll By: ANANDRISHIJI MEDICAL CENTREReport Date: 24-Jun-20234:57 PM

\* 0 6 2 3 2 4 0 1 2 \*

## **CBC-ESR**

	CBC-ES	SR	
Investigation	Result	Unit	Bio. Ref. Interval
HAEMOGLOBIN	14.1	g/dl	1318
TOTAL WBC COUNT	6800	/ cumm	4000-10000
RED BLOOD CELL COUNT	4.81	/cumm	4.32-5.72
WBC DIFFERENTIAL COUNT			
NEUTROPHILS	53	%	5070
LYMPHOCYTES	32	%	2040
EOSINOPHILS	06	%	06
MONOCYTES	09	%	0-10
BASOPHILS	00	%	01
RBC INDICES			
HEMATOCRIT	42.9	%	3754
MEAN CORPUSCULAR VOLUME	89.2	fl	78-92
MEAN CORPUSCULAR HEMOGLOBIN	29.3	pg	2832
MEAN CORPUSCULAR HEMOGLOBIN	32.9	g/dl	3237
CONCENTRATION			
RDW_CV	13.5	/ cumm	11.5-14.5
PLATELET COUNT	166000	/ cumm	150000-400000
MEAN PLATELET VOLUME	12.2	fl	7.4-10.4
PDW	16.3	fl	10-14
PCT	0.2	%	0.10-0.28
RED CELL DISTRIBUTION WIDTH (RDW-SD)	44.4	fl	
P-LCR	42.5	%	
PERIPHERAL BLOOD SMEAR			
ERYTHROCYTES	Normocytic	Normochromic	
	,		







Patient Name : MR. SANJEEV KUMAR

Age / Gender : 42 Years / Male

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### **CBC-ESR**

Investigation	Result	Unit	Bio. Ref. Interval			
LEUCOCYTES	Within nor	mal limits.				
THROMOBOCYTES	Adequate on smear					
ESR	02	mm/1hr.				
END OF REPORT						







Patient Name : MR. SANJEEV KUMAR

Age / Gender : 42 Years / Male

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 Patient ID
 : 062324012

Sample Coll By :ANANDRISHIJI MEDICAL CENTRE

Client Name : APOLLO

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\* 0 6 2 3 2 4 0 1 2 \*

### **TOTAL PSA**

Investigation	Result	Unit	Bio. Ref. Interval
TOTAL PSA	0.528	ng/ml	< 4.0 ng/mL

Specimen: Serum

Method: CLIA

#### Interpretation:

- .Increased levels are seen in prostatitis, benign hyperplasia, prostatic carcinoma, cirrhosis impotence, osteoporosis, pulmonary embolism, renal osteropathy, TUR and urinary retention.
- .Decreased levels are seen in castration, radiation therapy, prostatectomy, antiandrogen drugs (eg. finasteride).
- .PSA determination is used for monitoring of progress and efficiency of therapy in patients with prostate carcinoma or receiving hormonal therapy.
- .PSA has no circadian rhythm, but 6-7% variation can occur between specimens collected on same day.

Note: Clinical diagnosis should not be made on the findings of a single test result, but should integrate both clinical and laboratory data.







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

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Sample Coll By :ANANDRISHIJI MEDICAL CENTRE Report Date : 24-Jun-2023 4:30 PM

\* 0 6 2 3 2 4 0 1 2 \*

: APOLLO

# **Liver Function Test**

Investigation	Result	Unit	Bio. Ref. Interval	
ALKALINE PHOSPHATASE	115.5	U/L	53 - 128	
SGOT (AST)	30.1	U/L	0 -35	
SGPT (ALT)	44.0	U/L	0 - 45	
GGTP	38.9	U/L	0 - 55	
BILIRUBIN	1.00	mg/dL	0 - 1.2	
BILIRUBIN DIRECT	0.20	mg/dL	0 - 0.4	
BILIRUBIN INDIRECT	0.80	mg/dL	0 - 1.0	
TOTAL PROTEIN	6.7	g/dl	6.4 - 8.3	
ALBUMIN	4.1	gm/dl	3.5 - 5.2	
GLOBULIN	3	gm/dl	1.8 - 3.6	
A/G RATIO	1			
SGOT/SGPT RATIO	1	Ratio		







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

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# **GGT ( GAMMA GLUTAMYL TRANFERASE )**

Investigation	Result	Unit	Bio. Ref. Interval	
GGT (GAMMA GLUTAMYL TRANFERASE)	38.9	U/L	Male ≤ 55 Female ≤ 38	
METHOD	IFCC		Telliale 2 30	
SPECIMEN	Serum			
INSTRUMENT USED	Indiko			

### Interpretation:

The gamma-glutamyl transferase (GGT) test may be used to determine the cause of elevated alkaline phosphatase (ALP). Both ALP and GGT are elevated in disease of the bile ducts and in some liver diseases, but only ALP will be elevated in bone disease. Therefore, if the GGT level is normal in a person with a high ALP, the cause of the elevated ALP is most likely bone disease. An elevated GGT level suggests that something is damaging the liver. A low or normal GGT test result indicates that it is unlikely that a person has liver disease or has consumed any alcohol. A high GGT level can help rule out bone disease as the cause of an increased ALP level, but if GGT is low or normal, then an increased ALP is more likely due to bone disease.

Comment : Please correlate with clinical condition







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

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 : 24-Jun-2023
 4:55 PM

# **GLUCOSE - POST PRANDIAL(PP)**

Investigation	Result	Unit	Bio. Ref. Interval	
GLUCOSE - POST PRANDIAL(PP)				
GLUCOSE - POST PRANDIAL	107.0	mg/dL	70-140	

#### Interpretation:

A postprandial (PP) glucose test is a blood glucose test that determines the amount of a type of sugar, called glucose, in the blood after a meal. A 2-hour postprandial blood glucose test measures blood glucose exactly 2 hours after eating a meal, timed from the start of the meal. By this point blood sugar has usually gone back down in healthy people, but it may still be elevated in people with diabetes.

COMMENT Please correlate with clinical condition

TECHNOLOGY Spectrophotometry

NOTES Clinical diagnosis should not be made on the findings of a single

test result, but should integrate both clinical and laboratory

data.







**Patient Name Client Name** : MR. SANJEEV KUMAR : APOLLO

**Registration Date** : 24-Jun-2023 8:37 AM Age / Gender : 42 Years / Male

Ref. By Dr : APOLLO Sample Coll. Date : 24-Jun-2023 8:37 AM **Patient ID** : 062324012 **Authentication Date** : 24-Jun-2023 4:57 PM :ANANDRISHIJI MEDICAL CENTRE **Report Date** : 24-Jun-2023 4:30 PM

### LIPID PROFILE REPORT

Investigation	Result	Unit	Bio. Ref. Interval
TOTAL CHOLESTEROL	193.0	mg/dL	Desirable (< 200 )
			Borderline high (200 - 239 )
			High (> 240 )
HDL CHOLESTEROL - DIRECT	43.5	mg/dL	No Risk >60
			Moderate Risk 40 – 60
			High Risk <40
TRIGLYCERIDES	102.7	mg/dL	50-200
LDL CHOLESTEROL	129.0	mg/dL	Optimal (< 100 )
			Near optimal/above optimal
			(100-129)
			Borderline high (130-159)
			High (160-189)
			Very high (≥ 190)
VLDL CHOLESTEROL	20.5	mg/dL	5-40
TC/HDL CHOLESTEROL RATIO	4.4	Ratio	3.0-5.0
LDL / HDL RATIO	3.0	Ratio	1.5-3.5
NON HDL CHOLESTEROL	150	ng/ml	
HDL / LDL CHOLESTEROL RATIO	3	Ratio	1.5-3.5

#### Interpretation:

Sample Coll By

The lipid profile is used as part of a cardiac risk assessment to help determine an individual's risk of heart disease and to help make decisions about what treatment may be best if there is borderline or high risk. Lipids are a group of fats and fat-like substances that are important constituents of cells and sources of energy. A lipid profile typically includes: 1. Total cholesterol — this test measures all of the cholesterol in all the lipoprotein particles. 2. High-density lipoprotein cholesterol (HDL-C) — measures the cholesterol in HDL particles; often called "good cholesterol" because it removes excess cholesterol and carries it to the liver for removal. 3. Low-density lipoprotein cholesterol (LDL-C) — calculates the cholesterol in LDL particles; often called "bad cholesterol".

Comment : Please correlate with clinical condition







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

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## **HbA1C (GLYCOSYLATED HAEMOGLOBIN)**

Investigation	Value	Unit	
HBA1C (GLYCOSYLATED	5.4	%	Below 6.0 : Normal Value
HEMOGLOBIN), BLOOD			6.0-7.0 : Good Control
			7.0-8.0 : Fair Control
			8.0-10.0 : Unsatisfactory Control
			Above 10 : Poor Control
AVERAGE BLOOD GLUCOSE (ABG)	114.94	mg/dL	Below 136 : Normal Value
			137 - 172 : Good Control
			173 - 208 : Fair Control
			208 - 279 : Unsatisfactory Control
			Above 279: Poor Control

### **INTERPRETATION & REMARK**

### Interpretation

HbA1c is an indicator of glycemic control. HbA1c represents average glycemia over the past six to eight weeks. Glycation of hemoglobin occurs over the entire 120 day life span of the red blood cell, but with in this 120 days. Recent glycemia has the largest influence on the HbA1c value. Clinical studies suggest that a patient in stable control will have 50% of their HbA1c formed in the month before sampling, 25% in the month before that, and the remaining 25% in months two to four.

Comment Please correlate with with Clinical condition

Technology HPLC

Notes : Clinical diagnosis should not be made on the findings of a single test result, but should integrate both clinical and laboratory data.







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

 Ref. By Dr
 : APOLLO
 Sample Coll. Date
 : 24-Jun-2023
 8:37 AM

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 Authentication Date
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#### **THYROID FUNCTION TEST**

Investigation	Result	Unit	Bio. Ref. Interval	
TOTAL TRIIODOTHYRONINE (T3)	1.39	ng/ml	0.69-2.15	
TOTAL THYROXINE (T4)	9.60	ug/dl	5.2-12.7	
TSH	1.56	uIU/mL	0.3-4.5	

### T3/T4/TSH

Normal T3 concentrations do not necessarily reflect a normal – thyroid state. Certain thyroid disorders ( such as latent hypo – or hyperthyroidism , compensatory T3 over secretion in iodine deficiency , TBG over secretion) may also be associated with euthyroid T3 levels

In pregnancy , the Total T4 result may be incorrect , i.e., falsely –low .This assay should not be used as the only marker for thyroid disease evaluation during pregnancy. To ensure maximum diagnostic accuracy , thyroid status in pregnant women should be determined using thyroid function tests such as TSH , Free T4 , and clinical evaluation by the physician. Whether high or low , an abnormal TSH result indicates an excess or deficiency in the amount of thyroid hormone available to the body , but it does not indicate the reason . An abnormal TSH test result is usually followed by additional testing to investigate the cause of the increase or decrease.

Many medications – including aspirin and thyroid hormone replacement therapy – may affect thyroid gland function the result and their use should be discussed with the doctor prior to testing.

When a doctor adjusts a person's thyroid hormone replacement dosage, it is important to wait at least one to two months before checking the TSH again so that the new dose can have its full effect.

Extreme stress and acute illness may also affect TSH test result . Results may be low during the first trimester pregnancy. Serum TSH levels alone give no evidence of the presence or absence of thyroid disease. They must always be interpreted in context with the clinical picture and other diagnostic procedure.

A high TSH result often means an underactive thyroid gland that is not responding adequately to the stimulation of TSH due to some type of acute or chronic thyroid dysfunction. Rarely, a high TSH result can indicate a problem with the pituitary gland ,such as tumour producing unregulated levels of TSH.A high TSH can also occur when someone with a known thyroid disorder or who has their thyroid gland removed is receiving too little thyroid hormone medication. A low TSH result can indicate an overactive thyroid gland (hyperthyroidism) or excessive amounts of thyroid hormone medication in those who are being treated for an underactive (or removed) thyroid gland. Rarely, a low TSH result may indicate damage to the pituitary gland that prevents it from producing adequate amounts of TSH.

----- END OF REPORT







Age / Gender : 42 Years / Male Registration Date : 24-Jun-2023 8:37 AM

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### **RENAL FUNCTION TEST**

Result	Unit	Bio. Ref. Interval	
26.3	mg/dL	15-45	
0.75	mg/dL	0.5-1.5	
6.1	mg/dL	2.5-7.5	
138	mmol/L	136-146	
3.7	mmol/L	3.40-5.10	
101	mmol/L	98.0-106.0	
8.9	mg/dL	8.6 - 10.3	
	0.75 6.1 138 3.7 101	0.75 mg/dL 6.1 mg/dL 138 mmol/L 3.7 mmol/L 101 mmol/L	0.75 mg/dL 0.5-1.5 6.1 mg/dL 2.5-7.5 138 mmol/L 136-146 3.7 mmol/L 3.40-5.10 101 mmol/L 98.0-106.0

# Interpretation:

Renal function tests (RFT) are performed for evaluation of kidney function. The blood urea nitrogen or BUN test is primarily used, along with the creatinine test, to evaluate kidney function in a wide range of circumstances, to help diagnose kidney disease, and to monitor people with acute or chronic kidney dysfunction or failure. 1. Blood Urea Nitrogen (BUN) - Urea is a waste product formed in the liver when protein is metabolized. Urea is released by the liver into the blood and is carried to the kidneys, where it is filtered out of the blood and released into the urine. 2. Creatinine - Creatinine is a waste product produced by muscles from the breakdown of a compound called creatine. Almost all creatinine is filtered from the blood by the kidneys and released into the urine, so blood levels are usually a good indicator of how well the kidneys are working. 3. Uric acid - The uric acid blood test is used to detect high levels of this compound in the blood in order to help diagnose recurrent kidney stones and gout. The test is also used to monitor uric acid levels in people undergoing chemotherapy or radiation treatment for cancer.

Comment : Please correlate with clinical condition

Technology: Spectrophotometry

Notes : Clinical diagnosis should not be made on the findings of a single test result,

but should integrate both clinical and laboratory data.



