

CERTIFICATE OF MEDICAL FITNESS

0.4	
NAME: 18/1. Kellavathi. Gr.	-
AGE/ GENDER: 29 yu	
неіднт: 153 / ў.	WEIGHT: 51 Kg
IDENTIFICATION MARK:	_
BLOOD PRESSURE: 110/40 mm/14.	
PULSE: Fly wh	
CVS: Alormal	
1	
ANY OTHER DISEASE DIAGNOSED IN THE PAST:	
ALLERGIES, IF ANY:	
LIST OF PRESCRIBED MEDICINES:	
ANY OTHER REMARKS:	
I Certify that I have carefully examined Mr/Mrs. <u>Ka</u> of Mr <u>Granger ach</u> who has signed in r disease and is fit for employment.	y presence. He/ she has no physical
Signature of candidate	Dr. SATIS I KINI MD MEDICINE) Consultant Physician Signature of Medical Officer
Place: Spectrum diagnoshic the	altheane.
Date: 12 08 123	
Disclaimer: The patient has not been checked for COVID.	This certificate does not relate to the
"OVID STATUS OF the nationt evamined	

via status of the patient examinea











Dr. Ashok S Bsc., MBBS., D.O.M.S **Consultant Opthalmologist** KMC No: 31827

DATE: 12-68-23.

EYE EXAMINATIONP

NAME: Mg. Kalavalli	AGE: 27'Y	GENDER: F/M
	RIGHT EYE	LEFT EYE
Vision	6718:n	6)18:0B
Vision With glass		
Color Vision	Normal 	Normal
Anterior segment examination	Normal	Normal
Fundus Examination	Normal 	Normal
Any other abnormality	· Nill	Nill
Diagnosis/ impression	Normal	Normal
	B.\\$	Spedele OK SARODHE c., M.B.B.S., D.O.M.S. sultant & Surgeon productions





	Javil.	!	MRS KALAVATHI Female 29Years
			HR 78 bpm P 90 ms ORS 153 ms ORS 85 ms OT/OTc 392/448 ms P/QRS/T 72/69/-22 o RV5/SVI : 0:669/0.390 mV
No.	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\		Diagnosis Information: Sinus Arrhythmia T Wave Abnormality(III,aVF,V3,V6) Report Confirmed by: Report I Confirmed by:
			,V3,V6)



NAME	: MRS.KALAVATHI G	DATE :12/08/2023
AGE/SEX	: 29YEARS/FEMALE	REG NO:1208230068
REF BY	: APOLO CLINIC	

CHEST PA VIEW

Lung fields are clear.

Cardiovascular shadows are within normal limits.

Both CP angles are free.

Domes of diaphragm and bony thoracic cage are normal.

IMPRESSION: NORMAL CHEST RADIOGRAPH.

> DR.RAM PRAKASH G MDRD **CONSULTANT RADIOLOGIST**

RH1-19

Your suggestion / feedback is a valuable input for improving our services











PATIENT NAME	MRS KALAVATHI	ID NO	1208230068
AGE	29YEARS	SEX	FEMALE
REF BY	DR.APOLO CLINIC	DATE	12.08.2023

2D ECHO CARDIOGRAHIC STUDY

M-MODE

IVI-IVIOUL	
AORTA	21mm
LEFT ATRIUM	24mm
RIGHT VENTRICLE	18mm
LEFT VENTRICLE (DIASTOLE)	41mm
LEFT VENTRICLE(SYSTOLE)	32mm
VENTRICULAR SEPTUM (DIASTOLE)	07mm
VENTRICULAR SEPTUM (SYSTOLE)	08mm
POSTERIOR WALL (DIASTOLE)	08mm
POSTERIOR WALL (SYSTOLE)	09mm
FRACTIONAL SHORTENING	30%
EJECTION FRACTION	60%

DOPPLER /COLOUR FLOW

MITRAL VALVE	E-0.81 m/sec	A-0.46 m/sec	MILD MR
AORTIC VALVE	1.10m/sec		NO AR
PULMONARY VALVE			NO PR
	0.98 m/sec		
TRISCUSPID VALVE			*
		30mmHg	MILDTR











PATIENT NAME	MRS KALAVATHI	ID NO	1208230068
AGE	29YEARS	SEX	FEMALE
REF BY	DR.APOLO CLINIC	DATE	12.08.2023

2D ECHO CARDIOGRAHIC STUDY

LEFT VENTRICLE	SIZE& THICKNESS	NORMAL
CONTRACTILITY	REGIONAL GLOBAL	NO RWMA

RIGHT VENTRICLE : NORMAL
LEFT ATRIUM : NORMAL
RIGHT ATRIUM: NORMAL
MITRAL VALVE : NORMAL
AORTIC VALVE : NORMAL
PULMONARY VALVE: NORMAL
TRICUSPID VALVE: NORMAL
INTER ATRIAL SEPTUM :INTACT
INTER VENTRICULAR SEPTUM: INTACT
PERICARDIUM: NORMAL
OTHERS : - NIL

IMPRESSION

- > NO RWMA OF LV AT REST
- > NORMAL LV FUNCTION LVEF-60%
- > NORMAL CARDIAC CHAMBERS DIMENSIONS
- > MILD TR / MILD PAH
- > IAS & IVS INTACT
- > NORMAL IVC , NORMAL INSPIRATORY COLLAPSE
- > NO CLOT/ PERICARDIAL EFFUSION

ECHOTECHNICIAN

The science of radiology is based upon interpretation of shadows of normal and abnormal tissue. This is neither complete nor accurate; hence, findings should always be interpreted in to the light of clinico-pathological correction. This is a professional opinion









NAME AND LAB NO	MRS KALAVATHI G	Reg: 30068
AGE & SEX	29YRS	FEMALE
DATE AND AREA OF INTEREST	12.08.2023	ABDOMEN & PELVIS
REF BY	C/O APOLO CLINIC	

USG ABDOMEN AND PELVIS

LIVER:

Measures 13.1 cm. Normal in size with echotexture.

No e/o IHBR dilatation. No evidence of SOL.

Portal vein appears normal.

CBD appears normal. . No e/o calculus / SOL

GALL BLADDER:

Well distended. Wall appears normal. No e/o calculus/ neoplasm.

SPLEEN:

Measures 11.3 cm. Normal in size and echotexture. No e/o SOL/ calcification.

PANCREAS:

Normal in size and echotexture.

Pancreatic duct appears normal. No e/o calculus / calcifications.

RETROPERITONEUM:

Poor window.

RIGHT KIDNEY:

Measures 9.1 X4.0 cm. Right kidney is normal in size & echotexture

No evidence of calculus/ hydronephrosis.

LEFT KIDNEY:

Measures 10.8 X5.1 cm .Left kidney is normal in size & echotexture

No evidence of calculus/ hydronephrosis.

URETERS:

Bilateral ureters are not dilated.

URINARY BLADDER:

Minimally distended. No wall thickening/calculi.

UTERUS:

Anteverted, Normal in size and echotexture

Endometrium is normal.ET -10 mm.

OVARIES:

B/L ovaries normal in size and echotexture.

No evidence of ascites/pleural effusion.

IMPRESSION:

No significant sonological abnormality detected in the abdomen and pelvis.

DR AKSHATHA R BHAT

MDRD DNB FRCR









Age / Gender : 29 years / Female

: Dr. APOLO CLINIC Ref. By Dr.

: 1208230068 Reg. No. : Apollo Clinic $\mathbb{C}/0$

: 12-Aug-2023 09:45 AM **Bill Date**

Sample Col. Date: 12-Aug-2023 09:45 AM : 12-Aug-2023 02:55 PM **Result Date**

Report Status : Final

Test Name	Result	Unit	Reference Value	Method
Complete Haemogram-Whole B	lood EDTA			
Haemoglobin (HB)	12.0	g/dL	Female:12.0-15.0	Spectrophotmeter
Red Blood Cell (RBC)	4.46	million/cum	nm3.50 - 5.50	Volumetric Impedance
Packed Cell Volume (PCV)	36.6	%	Female: 36.0-45.0	Electronic Pulse
Mean corpuscular volume (MCV)	82.1	fL	78.0- 94.0	Calculated
Mean corpuscular hemoglobin (MCH)	26.5	pg	27.50-32.20	Calculated
Mean corpuscular hemoglobin concentration (MCHC)	32.3	%	33.00-35.50	Calculated
Red Blood Cell Distribution Width SD (RDW-SD)	34.3	fL	40.0-55.0	Volumetric Impedance
Red Blood Cell Distribution CV (RDW-CV)	15.3	%	Female: 12.20-16.10	Volumetric Impedance
Mean Platelet Volume (MPV)	11.0	fL	8.0-15.0	Volumetric Impedance
Platelet	2.2	lakh/cumm	1.50-4.50	Volumetric Impedance
Platelet Distribution Width (PDW)	24.6	%	8.30 - 56.60	Volumetric Impedance
White Blood cell Count (WBC)	6670.0	cells/cumm	Female: 4000.0-11000.0	Volumetric Impedance
Neutrophils	62.0	%	40.0-75.0	Light scattering/Manual
Lymphocytes	30.0	%	20.0-40.0	Light scattering/Manual
Eosinophils	2.0	%	0.0-6.0	Light scattering/Manual
Monocytes	5.0	%	0.0-8.0	Light scattering/Manual
Basophils	1.0	%	0.0-1.0	Light
Absolute Neutrophil Count	4.02	10^3/uL	2.0- 7.0	scattering/Manual Calculated

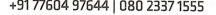
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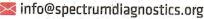
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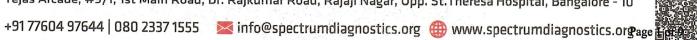


















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Test Name	Result	Unit	Reference Value	Method
Absolute Lymphocyte Count	2.01	10^3/uL	1.0-3.0	Calculated
Absolute Monocyte Count	0.52	10^3/uL	0.20-1.00	Calculated
Absolute Eosinophil Count	110	cells/cumm	40-440	Calculated
Absolute Basophil Count	0.01	10^3/uL	0.0-0.10	Calculated
Erythrocyte Sedimentation Rate (ESR)	45	mm/hr	Female: 0.0-20.0	Westergren

1208230068

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Peripheral Smear Examination-Whole Blood EDTA

Method: (Microscopy-Manual)

RBC'S : Normocytic Normochromic.

WBC'S : Are normal in total number, morphology and distribution.

Platelets : Adequate in number and normal in morphology.

No abnormal cells or hemoparasites are present.

Impression: Normocytic Normochromic Blood picture.

Blood Group & Rh Typing-Whole Blood EDTA

Blood Group Slide/Tube

agglutination

Rh Type **Positive** Slide/Tube

agglutination

Note: Confirm by tube or gel method.

Comments: ABO blood group system, the classification of human blood based on the inherited properties of red blood cells (erythrocytes) as determined by the presence or absence of the antigens A and B, which are carried on the surface of the red cells. Persons may thus have type A, type B, type O, or type AB blood.



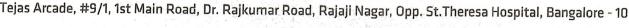
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Printed On : 12 Aug, 2023 07:26 pm

Dr. Nithun Reddy C,MD,Consultant Pathologist











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Test Name	Result	Unit	Reference Value	Method
Fasting Blood Sugar (FBS)-	78	mg/dL	60.0-110.0	Hexo Kinase

1208230068

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Comments: Glucose, also called dextrose, one of a group of carbohydrates known as simple sugars (monosaccharides). Glucose has the molecular formula C₆H₁₂O₆. It is found in fruits and honey and is the major free sugar circulating in the blood of higher animals. It is the source of energy in cell function, and the regulation of its metabolism is of great importance (fermentation; gluconeogenesis). Molecules of starch, the major energy-reserve carbohydrate of plants, consist of thousands of linear glucose units. Another major compound composed of glucose is cellulose, which is also linear. Dextrose is the molecule D-glucose. Blood sugar, or glucose, is the main sugar found in the blood. It comes from the food you eat, and it is body's main source of energy. The blood carries glucose to all of the body's cells to use for energy. Diabetes is a disease in which your blood sugar levels are too high.Usage: Glucose determinations are useful in the detection and management of Diabetes mellitus.

Note: Additional tests available for Diabetic control are Glycated Hemoglobin (HbA1c), Fructosamine & Microalbumin urine

Comments: Conditions which can lead to lower postprandial glucose levels as compared to fasting glucose are excessive insulin release, rapid gastric emptying & brisk glucose absorption.

Probable causes: Early Type II Diabetes / Glucose intolerance, Drugs like Salicylates, Beta blockers, Pentamidine etc., Alcohol , Dietary - Intake of excessive carbohydrates and foods with high glycemic index? Exercise in between samples? Family history of Diabetes, Idiopathic, Partial / Total Gastrectomy.



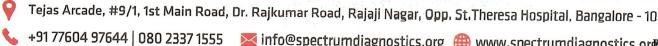
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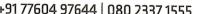
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Test Name	Result	Unit	Reference Value	Method
Post Prandial Urine Sugar	Negative		Negative	Dipstick/Benedicts(Man
Post prandial Blood Glucose (PPBS)-Plasma	94	mg/dL	80.0-150.0	Hexo Kinase

1208230068

: 1208230068

Comments: Glucose, also called dextrose, one of a group of carbohydrates known as simple sugars (monosaccharides). Glucose has the molecular formula $C_6H_{12}O_6$. It is found in fruits and honey and is the major free sugar circulating in the blood of higher animals. It is the source of energy in cell function, and the regulation of its metabolism is of great importance (fermentation; gluconeogenesis). Molecules of starch, the major energy-reserve carbohydrate of plants, consist of thousands of linear glucose units. Another major compound composed of glucose is cellulose, which is also linear. Dextrose is the molecule D-glucose. Blood sugar, or glucose, is the main sugar found in the blood. It comes from the food you eat, and it is body's main source of energy. The blood carries glucose to all of the body's cells to use for energy. Diabetes is a disease in which your blood sugar levels are too high.Usage: Glucose determinations are useful in the detection and management of Diabetes mellitus.

Note: Additional tests available for Diabetic control are Glycated Hemoglobin (HbA1c), Fructosamine & Microalbumin urine

UHID

Comments: Conditions which can lead to lower postprandial glucose levels as compared to fasting glucose are excessive insulin release, rapid gastric emptying & brisk glucose absorption.

Probable causes: Early Type II Diabetes / Glucose intolerance, Drugs like Salicylates, Beta blockers, Pentamidine etc., Alcohol, Dietary - Intake of excessive carbohydrates and foods with high glycemic index? Exercise in between samples? Family history of Diabetes, Idiopathic, Partial / Total Gastrectomy.



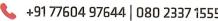
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SCAN FOR LOCATION





Tejas Arcade, #9/1, 1st Main Road, Dr. Rajkumar Road, Rajaji Nagar, Opp. St.Theresa Hospital, Bangalore - 10







Age / Gender : 29 years / Female

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Test Name	Result	Unit	Reference Value	Method
Thyroid function tests (TF Serum	Γ)-			
Tri-Iodo Thyronine (T3)-So	erum 0.78	ng/mL	0.60-1.81	Chemiluminescence Immunoassay (CLIA)
Thyroxine (T4)-Serum	7.2	μg/dL	5.50-12.10	Chemiluminescence Immunoassay (CLIA)
Thyroid Stimulating Hormo (TSH)-Serum	one 1.61	μIU/mL	0.35-5.50	Chemiluminescence Immunoassay (CLIA)

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Comments: Triiodothyronine (T3) assay is a useful test for hyperthyroidism in patients with low TSH and normal T4 levels. It is also used for the diagnosis of T3 toxicosis. It is not a reliable marker for Hypothyroidism. This test is not recommended for general screening of the population without a clinical suspicion of hyperthyroidism.

Reference range: Cord: (37 Weeks): 0.5-1.41, Children:1-3 Days: 1.0-7.40,1-11 Months: 1.05-2.45,1-5 Years: 1.05-2.69,6-10 Years: 0.94-2.41,11-15 Years: 0.82-2.13, Adolescents (16-20 Years): 0.80-2.10

Reference range: Adults: 20-50 Years: 0.70-2.04, 50-90 Years: 0.40-1.81,

Reference range in Pregnancy: First Trimester: 0.81-1.90, Second Trimester: 1.0-2.60

Increased Levels: Pregnancy, Graves disease, T3 thyrotoxicosis, TSH dependent Hyperthyroidism, increased Thyroid-binding globulin (TBG). Decreased Levels: Nonthyroidal illness, hypothyroidism, nutritional deficiency, systemic illness, decreased Thyroid-binding globulin (TBG).

Comments: Total T4 levels offer a good index of thyroid function when TBG is normal and non-thyroidal illness is not present. This assay is useful for monitoring treatment with synthetic hormones (synthetic T3 will cause low total T4). It also helps to monitor treatment of Hyperthyroidism with Thiouracil or other anti-thyroid drugs.

Reference Range: Males: 4.6-10.5, Females: 5.5-11.0, 60 Years: 5.0-10.70, Cord: 7.40-13.10, Children: 1-3 Days: 11.80-22.60, 1-2 Weeks: 9.90-16.60,1-4 Months: 7.20-14.40,1-5 Years: 7.30-15.0,5-10 Years: 6.4-13.3

1-15 Years: 5.60-11.70, Newborn Screen: 1-5 Days: >7.5,6 Days :>6.5

Increased Levels: Hyperthyroidism, increased TBG, familial dysalbuminemic hyperthyroxinemia, Increased transthyretin, estrogen therapy, pregnancy. Decreased Levels: Primary hypothyroidism, pituitary TSH deficiency, hypothalamic TRH deficiency, non thyroidal illness, decreased TBG.

Comments: TSH is a glycoprotein hormone secreted by the anterior pituitary. TSH is a labile hormone & is secreted in a pulsatile manner throughout the day and is subject to several non-thyroidal pituitary influences. Significant variations in TSH can occur with circadian rhythm, hormonal status, stress, sleep deprivation, caloric intake, medication & circulating antibodies. It is important to confirm any TSH abnormality in a fresh specimen drawn after ~ 3 weeks before assigning a diagnosis, as the cause of an isolated TSH abnormality.

Reference range in Pregnancy: I- trimester:0.1-2.5; II -trimester:0.2-3.0; III- trimester:0.3-3.0

Reference range in Newborns: 0-4 days: 1.0-39.0; 2-20 Weeks:1.7-9.1

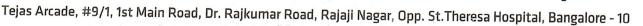
Increased Levels: Primary hypothyroidism, Subclinical hypothyroidism, TSH dependent Hyperthyroidism and Thyroid hormone resistance.

Decreased Levels: Graves disease, Autonomous thyroid hormone secretion, TSH deficiency.

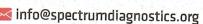
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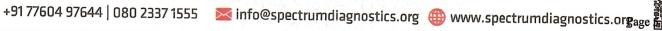
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Test Name	Result	Unit	Reference Value	Method
Gamma-Glutamyl Transferase (GGT)-Serum	28.00	U/L	Female: 5.0-55.0	Other g-Glut- 3-carboxy-4 nitro

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Comments: Gamma-glutamyltransferase (GGT) is primarily present in kidney, liver, and pancreatic cells. Small amounts are present in other tissues. Even though renal tissue has the highest level of GGT, the enzyme present in the serum appears to originate primarily from the hepatobiliary system, and GGT activity is elevated in any and all forms of liver disease. It is highest in cases of intra- or posthepatic biliary obstruction, reaching levels some 5 to 30 times normal. GGT is more sensitive than alkaline phosphatase (ALP), leucine aminopeptidase, aspartate transaminase, and alanine aminotransferase in detecting obstructive jaundice, cholangitis, and cholecystitis; its rise occurs earlier than with these other enzymes and persists longer. Only modest elevations (2-5 times normal) occur in infectious hepatitis, and in this condition, GGT determinations are less useful diagnostically than are measurements of the transaminases. High elevations of GGT are also observed in patients with either primary or secondary (metastatic) neoplasms. Elevated levels of GGT are noted not only in the sera of patients with alcoholic cirrhosis but also in the majority of sera from persons who are heavy drinkers. Studies have emphasized the value of serum GGT levels in detecting alcohol-induced liver disease. Elevated serum values are also seen in patients receiving drugs such as phenytoin and phenobarbital, and this is thought to reflect induction of new enzyme activity.

RFT (Urea, Creatinine, BUN, Na+, K+, Cl-, RBS Uric acid, HB)

RFT (Renal Function Test)-

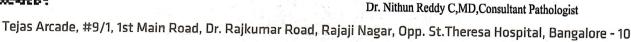
Serum

Serum				
Urea-Serum	15.8	mg/dL	Female: 06 - 40	Urease
Creatinine-Serum	0.73	mg/dL	Female: 0.5 - 1.1	Modified kinetic Jaffe
Blood Urea Nitrogen (BUN)- Serum	7.4	mg/dL	Female: 6 - 20	:GLDH,Kinetic Assay
Sodium (Na+)-Serum	141.8	mmol/L	Female: 135 - 145	ISE-Direct
Potassium (K+)-Serum	3.87	mmol/L	Female: 3.5 - 5.5	ISE-Direct
Chloride (Cl-)-Serum Uric Acid-Serum	103.20 3.10	mmol/L mg/dL	94.0 - 110.0 Female: 2.60 - 6.00	ISE-Direct Uricase PAP

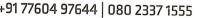


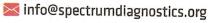
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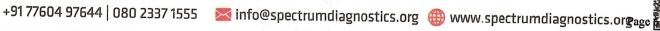


















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Test Name	Result	Unit	Reference Value	Method
Lipid Profile-Serum				
Cholesterol Total-Serum	152.00	mg/dL	0.0-200	Cholesterol Oxidase/Peroxidase
Triglycerides-Serum	146.00	mg/dL	0.0-150	Lipase/Glycerol Dehydrogenase
High-density lipoprotein (HDL) Cholesterol-Serum	33.00	mg/dL	40.0-60.0	Accelerator/Selective Detergent
Non-HDL cholesterol-Serum	119	mg/dL	0.0-130	Calculated
Low-density lipoprotein (LDL) Cholesterol-Serum	90	mg/dL	0.0-100.0	Cholesterol esterase and cholesterol oxidase
Very-low-density lipoprotein (VLDL) cholesterol-Serum	29	mg/dL	0.0-40	Calculated
Cholesterol/HDL Ratio-Serum	4.61	Ratio	0.0-5.0	Calculated

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Interpretation:

Parameter	Desirable	Borderline High	High	Very High
Total Cholesterol	<200	200-239	>240	
Triglycerides	<150	150-199	200-499	>500
Non-HDL cholesterol	<130	160-189	190-219	>220
Low-density lipoprotein (LDL) Cholesterol	<100	100-129	160-189	>190

Comments: As per Lipid Association of India (LAI), for routine screening, overnight fasting preferred but not mandatory. Indians are at very high risk of developing Atherosclerotic Cardiovascular (ASCVD). Among the various risk factors for ASCVD such as dyslipidemia, Diabetes Mellitus, sedentary lifestyle, Hypertension, smoking etc., dyslipidemia has the highest population attributable risk for MI both because of direct association with disease pathogenesis and very high prevalence in Indian population. Hence monitoring lipid profile regularly for effective management of dyslipidemia remains one of the most important healthcare targets for prevention of ASCVD. In addition, estimation of ASCVD risk is an essential, initial step in the management of individuals requiring primary prevention of ASCVD. In the context of lipid management, such a risk estimate forms the basis for several key therapeutic decisions, such as the need for and aggressiveness of statin therapy.



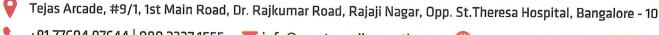
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Age / Gender : 29 years / Female

: Dr. APOLO CLINIC

Reg. No. : 1208230068

Ref. By Dr.

C/o : Apollo Clinic **Bill Date** : 12-Aug-2023 09:45 AM

Sample Col. Date: 12-Aug-2023 09:45 AM : 12-Aug-2023 02:55 PM **Result Date**

Report Status : Final

Test Name	Result	Unit	Reference Value	Method
Glycosylated Haemoglobin (HbA1c)-Whole Blood EDTA				
Glycosylated Haemoglobin (HbA1c)	5.30	%	Non diabetic adults:<5.7 At risk (Prediabetes): 5.7 - 6.4 Diagnosing Diabetes:>= 6.5 Diabetes Excellent Control: 6-7 Fair to good Control: 7-8	HPLC
			Unsatisfactory Control :8-10 Poor Control :>10	
Estimated Average	105.41	mg/dL		Calculated

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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.

Comments: 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 as compared to blood and urinary glucose determinations.



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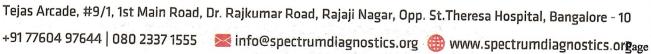
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: Final Report Status

Test Name	Result	Unit	Reference Value	Method
Urine Routine Examination-	-Urine			
Physical Examination				
Colour	Pale Yellow		Pale Yellow	Visual
Appearance	Clear		Clear	Visual
Reaction (pH)	5.5		5.0 - 7.5	Dipstick
Specific Gravity	1.020		1.000 - 1.030	Dipstick
Biochemical Examination				
Albumin	Negative		Negative	Dipstick/Precipitation
Glucose	Negative		Negative	Dipstick/Benedicts
Bilirubin	Negative		Negative	Dipstick/Fouchets
Ketone Bodies	Negative		Negative	Dipstick/Rotheras
Urobilinogen	Normal		Normal	Dipstick/Ehrlichs
Nitrite	Negative		Negative	Dipstick
Microscopic Examination	1 (oguil vo			
Pus Cells	2-3	hpf	0.0 - 5.0	Microscopy
Epithelial Cells	2-3	hpf	0.0 - 10.0	Microscopy
RBCs	Absent	hpf	Absent	Microscopy
Casts	Absent	mp.	Absent	Microscopy
Crystals	Absent		Absent	Microscopy
Others	Absent		Absent	Microscopy

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Comments: The kidneys help infiltration of the blood by eliminating waste out of the body through urine. They also regulate water in the body by conserving electrolytes, proteins, and other compounds. But due to some conditions and abnormalities in kidney function, the urine may encompass some abnormal constituents, which are not normally present. A complete urine examination helps in detecting such abnormal constituents in urine. Several disorders can be detected by identifying and measuring the levels of such substances. Blood cells, bilirubin, bacteria, pus cells, epithelial cells may be present in urine due to kidney disease or infection. Routine urine examination helps to diagnose kidney diseases, urinary tract infections, diabetes and other metabolic disorders.



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Dr. Nithun Reddy C,MD,Consultant Pathologist





