

CERTIFICATE OF MEDICAL FITNESS

NAME: Sakaray Ramish.
AGE/ GENDER: 1114
HEIGHT: 174 Cron. WEIGHT: 90-1 kg
IDENTIFICATION MARK:
BLOOD PRESSURE: 140/96 roomtig.
PULSE: 81 Mim
CVS:
RS:P (Noomax.
ANY OTHER DISEASE DIAGNOSED IN THE PAST: NIL
ALLERGIES, IF ANY:
LIST OF PRESCRIBED MEDICINES:
ANY OTHER REMARKS:
of Ms Sheshappa who has signed in my presence. He/ she has no physical
disease and is fit for employment.
Reg. No. 62806
Signature of candidate Signature of Medical Officer
Place: Spectoum Diagnostic & health core
Date: 30 04 24

Disclaimer: The patient has not been checked for COVID. This certificate does not relate to the covid status of the patient examined





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

DATE: 2004.24.

EYE EXAMINATION

NAME: Mg. Sa.	Kosey
---------------	-------

AGE: 41 Y

GENDER: F/M

RIGHT EYE

LEFT EYE

Vision

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

Consultant (Opthalmologist)





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					2 1 2 2	2					EAMESH	
							P/QRS/T RV5/SV1	QT/QTc	בי מ	j 'T	ţ	20-04-2024
			\$ \$ \$ }	1			: 49/24/41 ° : 1.019/0.960 mV	353/409 ms	: 169 ms		; 80 opm	09:4
	\$		\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	\{ \(\)	VS	<u>X</u>	v V Report	200	Prolonged	Sinus	Diagnos	FORBPL
							Confirmed by:	IIIIai V wave(III)			Diagnosis Information:	
\\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\			<u>} </u>	<u>}</u>	<u> </u>							
			} }	} }	<u>} </u>	HPI-MANAGED ROSS	CONTRACTOR OF THE PARTY OF THE					
) 	<u> </u>	<u>}</u>	<u> </u>	744 T		43/16	y			



SPECTRUM DIAGNOSTICS

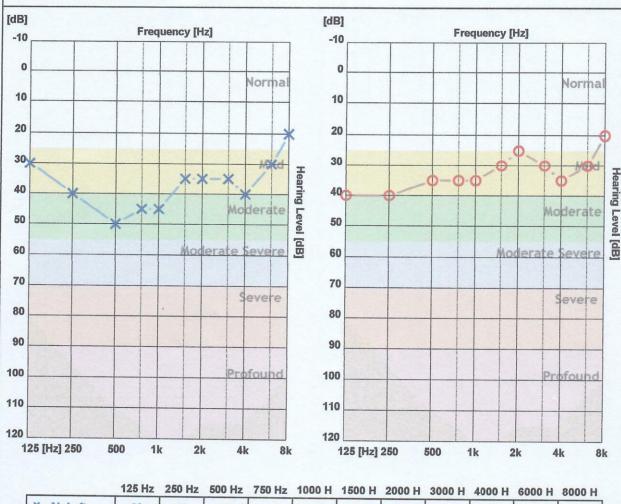
Bangalore

Patient ID: 0325

Name: SAKARAY RAMESH CR Number: 20240420093713 Registration Date: 20-Apr-2024 Age: 41

Gender : Male

Operator: spectrum diagnostics



	125 Hz	250 Hz	500 Hz	750 Hz	1000 H	1500 H	2000 H	3000 H	4000 H	6000 H	8000 H
X - Air Left	30	40	50	45	45	35	35	35	40	30	20
O - Air Right	40	40	35	35	35	30	25	30	35	30	20
> - Bone Left											
< - Bone Right							-				

	Average	High	Mid	Low
AIR Left	36.82 dB	31.25 dB	38.33 dB	41.25 dB
AIR Right	32.27 dB	28.75 dB	30.00 dB	37.50 dB

Clinical Notes:

Not Found







MESH DATE : 20/04/2024
REG NO :2004240021
11011011210021

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 RIKHIT MAGANLAL CONSULTANT RADIOLOGIST

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





PATIENT NAME	MR SAKARAY RAMESH	ID NO	2004240021
AGE	41YEARS	SEX	MALE
REFBY	DR.APOLO CLINIC	DATE	20.04.2024

2D ECHO CARDIOGRAHIC STUDY

IVI	WODE
AORTA	35mm
LEFT ATRIUM	38mm
RIGHT VENTRICLE	20mm
LEFT VENTRICLE (DIASTOLE)	46mm
LEFT VENTRICLE(SYSTOLE)	31mm
VENTRICULAR SEPTUM (DIASTOLE)	12mm
VENTRICULAR SEPTUM (SYSTOLE)	13mm
POSTERIOR WALL (DIASTOLE)	12mm
POSTERIOR WALL (SYSTOLE)	11mm
FRACTIONAL SHORTENING	30%
EJECTION FRACTION	60%

DOPPLER /COLOUR FLOW

Mitral Valve Velocity: MVE-0.74m/s MVA - 0.51m/s E/A-1.46

Tissue Doppler: e' (Septal) -8cm/s E/e'(Septal) -9

Velocity/ Gradient across the Pulmonic valve : 0.83m/s 3mmHg

Max. Velocity / Gradient across the Aortic valve: 1.19m/s 4mmHg

Velocity / Gradient across the Tricuspid valve : 1.87 m/s 19mmHg







PATIENT NAME	MR SAKARAY RAMESH	ID NO	2004240021
AGE	41YEARS	SEX	MALE
REF BY	DR.APOLO CLINIC	DATE	20.04.2024

2D ECHO CARDIOGRAHIC STUDY

LEFT VENTRICLE	SIZE& THICKNESS	LVH
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	577	NORMAL	
INTER ATRIAL SEPTUM	:	INTACT	
INTER VENTRICULAR SEPT	UM:	INTACT	3277.0
PERICARDIUM	;	NORMAL	
OTHERS	:	- NIL	

IMPRESSION

- NO REGIONAL WALL MOTION ABNORMALITY PRESENT
- NORMAL VALVES AND DIMENSIONS
- NORMAL LV FUNCTION, LVEF- 60%
- LEFT VENTRICULAR HYPERTROPHY
- TRIVIAL TR / NO PAH
- NORMAL RV FUNCTION
- NO CLOT / VEGETATION / EFFUSION



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.





NAME AND LAB NO	MR SAKARAY RAMESH	REG-40021
AGE & SEX	41 YRS	MALE
DATE AND AREA OF INTEREST	20.04.2024	ABDOMEN & PELVIS
REF BY	C/O APOLO CLINIC	ADDOMEN & PELVIS

USG ABDOMEN AND PELVIS

LIVER:

Measures 15.0 cm. Normal in size and shows diffuse increased echogenicity

No e/o IHBR dilatation. No evidence of focal lesion. Portal vein appears normal. CBD appears normal.

GALL BLADDER:

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

SPLEEN:

Normal in size and echotexture. No e/o focal lesion.

PANCREAS:

Head and body appears normal . Tail obscured by bowel gas shadows .

RETROPERITONEUM:

Suboptimal visualised due to bowel gas

RIGHT KIDNEY:

Right kidney ,is normal in size & echotexture.

No evidence of calculus/ hydronephrosis.

No solid lesions.

LEFT KIDNEY:

Left kidney ,is normal in size & echotexture.

No evidence of calculus/ hydronephrosis.

No solid lesions.

URINARY BLADDER:

Well distended. No wall thickening/ calculi.

PROSTATE:

Normal in size and echotexture.

No evidence of ascites/pleural effusion.

IMPRESSION:

Grade I fatty liver.

Suggested clinical / lab correlation

DR PURNIMA PUJAR MDRD **CONSULTANT RADIOLOGIST**









Age / Gender : 41 years / Male

Ref. By Dr. : Dr. APOLO CLINIC

Reg. No. : 2004240021

C/o

Bill Date : 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM **Result Date** : 20-Apr-2024 12:52 PM

Report Status : Final

2004240021 : Apollo Clinic

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Test Name Result Unit Reference Value Method Fasting Blood Sugar (FBS)-93 mg/dL 60.0-110.0 Hexo Kinase Plasma

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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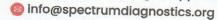
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Age / Gender : 41 years / Male

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Test Name	Result	Unit	Reference Value	Method
Glycosylated Haemoglobin (HbA1c)-Whole Blood EDTA	A			
Glycosylated Haemoglobin	4.70	%	Non diabetic adults :<5.7	HPLC
(HbA1c)			At risk (Prediabetes): 5.7 - 6.4	
			Diagnosing Diabetes :>= 6.5	
			Diabetes	
			Excellent Control: 6-7	
			Fair to good Control: 7-8 Unsatisfactory Control: 8-10	
Estimated Assessed	00.40		Poor Control :>10	
Estimated Average Glucose(eAG)	88.18	mg/dL		Calculated

: 2004240021

2004240021

UHID

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

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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Test Name	Result	Unit	Reference Value	Method
Thyroid function tests (TF) Serum	Γ)-			
Tri-Iodo Thyronine (T3)-So	erum 1.00	ng/mL	Male: 0.60 - 1.81	Chemiluminescence Immunoassay
Thyroxine (T4)-Serum	7.80	μg/dL	Male: 5.50 - 12.10	(CLIA) Chemiluminescence Immunoassay
Thyroid Stimulating Hormo (TSH)-Serum	one 3.35	μIU/mL	Male: 0.35 - 5.50	(CLIA) Chemiluminescence Immunoassay (CLIA)

2004240021

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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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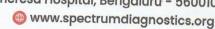
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Test Name	Result	Unit	Reference Value	Method
LFT-Liver Function Test -Seru	m	411 Hall (1997)		
Bilirubin Total-Serum	0.91	mg/dL	0.2-1.0	Caffeine Benzoate
Bilirubin Direct-Serum	0.15	mg/dL	0.0-0.2	Diazotised Sulphanilic Acid
Bilirubin Indirect-Serum	0.76	mg/dL	Male: 0.0 - 1.10	Direct Measure
Aspartate Aminotransferase (AST/SGOT)-Serum	22.00	U/L	Male: 15.0 - 37.0	UV with Pyridoxal - 5 -
Alanine Aminotransferase (ALT/SGPT)-Serum	28.00	U/L	Male: 16.0 - 63.0	Phosphate UV with Pyridoxal - 5 -
Alkaline Phosphatase (ALP)- Serum	49.00	U/L	Male: 45.0 - 117.0	Phosphate PNPP,AMP- Buffer
Protein, Total-Serum	6.78	g/dL	6.40-8.20	Biuret/Endpoint- With Blank
Albumin-Serum	4.51	g/dL	Male: 3.40 - 5.50	Bromocresol Purple
Globulin-Serum	2.27	g/dL	2.0-3.50	Calculated
Albumin/Globulin Ratio-Serun	1.99	Ratio	0.80-2.0	Calculated

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SCAN FOR LOCATION







Age / Gender : 41 years / Male Ref. By Dr.

: Dr. APOLO CLINIC Reg. No. : 2004240021

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Sample Col. Date: 20-Apr-2024 08:51 AM **Result Date** : 20-Apr-2024 12:52 PM

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Test Name	Result	Unit	Reference Value	Method
Lipid Profile-Serum				
Cholesterol Total-Serum	173.00	mg/dL	Male: 0.0 - 200	Cholesterol
Triglycerides-Serum	205.00	mg/dL	Male: 0.0 - 150	Oxidase/Peroxidase Lipase/Glycerol
High-density lipoprotein (HDL) Cholesterol-Serum	48.00	mg/dL	Male: 40.0 - 60.0	Dehydrogenase Accelerator/Selective
Non-HDL cholesterol-Serum	125	mg/dL	Male: 0.0 - 130	Detergent
Low-density lipoprotein (LDL) Cholesterol-Serum	57075550	mg/dL	Male: 0.0 - 100.0	Calculated Cholesterol esterase and cholesterol
Very-low-density lipoprotein (VLDL) cholesterol-Serum	41	mg/dL	Male: 0.0 - 40	oxidase Calculated
Cholesterol/HDL Ratio-Serum	3.60	Ratio	Male: 0.0 - 5.0	Calculated
nterpretation.				

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Interpretation:

Desirable	Borderline High	High	Very High
<200	200-239		very rigi
<150	150-199		>500
<130	160-189		>220
<100			>190
	<200 <150 <130	<200 200-239 <150 150-199 <130 160-189	<200

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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Test Name	Result	Unit	Reference Value	Method
Calcium, Total- Serum	9.00	mg/dL	8.50-10.10	Spectrophotometry (O- Cresolphthalein
Gamma-Glutamyl Transferase (GGT)-Serum	16.00	U/L	Male: 15.0-85.0	complexone) Other g-Glut-3-
			Female: 5.0-55.0	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.



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Test Name	Result	Unit	Reference Value	Method	
Prostate-Specific Antig	gen(PSA)-0.61	ng/mL	0.0-4.0	CLIA	

2004240021

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Note: 1. This is a recommended test for detection of prostate cancer along with Digital Rectal Examination (DRE) in males above 50 years of age.

2. False negative / positive results are observed in patients receiving mouse monoclonal antibodies for diagnosis or therapy.

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3. PSA levels may appear consistently elevated / depressed due to the interference by heterophilic antibodies & nonspecific protein binding.

4. Immediate PSA testing following digital rectal examination, ejaculation, prostatic massage, indwelling catheterization, ultrasonography and needle biopsy of prostate is not recommended as they falsely elevate levels

5. PSA values regardless of levels should not be interpreted as absolute evidence of the presence or absence of disease. All values should be

clinical findings and results of other investigations

6. Sites of Non-prostatic PSA production are breast epithelium, salivary glands, periurethral & anal glands, cells of male urethra & breast milk

7. Physiological decrease in PSA level by 18% has been observed in hospitalized /sedentary patients either due to supine position or suspended sexual activity.

Recommended Testing Intervals: Pre-operatively (Baseline), 2-4 days post-operatively, Prior to discharge from hospital, Monthly followup if levels are high or show a rising trend.

Clinical Use: -An aid in the early detection of Prostate cancer when used in conjunction with Digital rectal examination in males more than 50 years of age and in those with two or more affected first degree relatives.

-Followup and management of Prostate cancer patients

-Detect metastatic or persistent disease in patients following surgical or medical treatment of Prostate cancer. Increased Levels: Prostate cancer, Benign Prostatic Hyperplasia, Prostatitis, Genitourinary infections.



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Test Name Result Unit Reference Value Method

Fasting Urine Glucose-Urine Negative Negative Dipstick/Benedicts (Manual)



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Test Name	Result	Unit	Reference Value	Method
Urine Routine Examination-	Urine			
Physical Examination				
Colour	Pale Yellow		Pale Yellow	V:1
Appearance	Clear		Clear	Visual
Reaction (pH)	5.5		5.0-7.5	Visual
Specific Gravity	1.025		1.000-1.030	Dipstick
Biochemical Examination			1.000-1.050	Dipstick
Albumin	Negative		Negative	Di
Glucose	Negative		Negative	Dipstick/Precipitation
Bilirubin	Negative		Negative	Dipstick/Benedicts
Ketone Bodies	Negative			Dipstick/Fouchets
Urobilinogen	Normal		Negative Normal	Dipstick/Rotheras
Nitrite	Negative			Dipstick/Ehrlichs
Microscopic Examination	riogative		Negative	Dipstick
Pus Cells	2-4	h.f	0.0.5.0	
Epithelial Cells	1-2	hpf	0.0-5.0	Microscopy
RBCs	2-3	hpf	0.0-10.0	Microscopy
Casts	Absent	hpf	Absent	Microscopy
Crystals			Absent	Microscopy
or y seeing	Crystal Proger		Absent	Microscopy
Others	Crystal Preser Absent	ıı	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 byidentifying 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,



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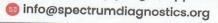
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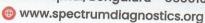
Dr. Nithun Reddy C,MD,Consultant Pathologist

Tejas Arcade, #9/1, 1st Main Road, Dr. Rajkumar Road, Rajajinagar, Opp. St. Theresa Hospital, Bengaluru - 560010













Age / Gender : 41 years / Male

Ref. By Dr. : Dr. APOLO CLINIC

Reg. No. : 2004240021

C/o : Apollo Clinic **Bill Date** : 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM

: 20-Apr-2024 01:11 PM Report Status : Final

Result Date

Test Name Result Unit Reference Value Method Post prandial Blood Glucose 103 mg/dL 70-140 Hexo Kinase (PPBS)-Plasma

2004240021

: 2004240021

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

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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



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C/o : Apollo Clinic **Bill Date**

: 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM

Result Date : 20-Apr-2024 01:11 PM Report Status : Final

Test Name	Result	Unit	Reference Value	Method	
KFT (Kidney Function Test)	:				**************************************
Blood Urea Nitrogen (BUN)- Serum	15.50	mg/dL	7.0-18.0	GLDH,Kinetic Assay	
Creatinine-Serum	0.82	mg/dL	Male: 0.70-1.30 Female: 0.55-1.02	Modified	
Uric Acid-Serum	7.08	mg/dL	Male: 3.50-7.20 Female: 2.60-6.00	kinetic Jaffe Uricase PAP	
Sodium (Na+)-Serum	139.3	mmol/L	135.0-145.0	Ion-Selective Electrodes (ISE)	
Potassium (K+)-Serum	4.06	mmol/L	3.5 to 5.5	Ion-Selective Electrodes (ISE)	,
Chloride(Cl-)-Serum	107.10	mmol/L	96.0-108.0	Ion-Selective Electrodes (ISE)	

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Comments: Renal Function Test (RFT), also called kidney function tests, are a group of tests performed to evaluate the functions of the kidneys. The kidneys play a vital role in removing waste, toxins, and extra water from the body. They are responsible for maintaining a healthy balance of water, salts, and minerals such as calcium, sodium, potassium, and phosphorus. They are also essential for blood pressure control, maintenance of the body's pH balance, making red blood cell production hormones, and promoting bone health. Hence, keeping your kidneys healthy is essential for maintaining overall health. It helps diagnose inflammation, infection or damage in the kidneys. The test measures Uric Acid, Creatinine, BUN and electrolytes in the blood to determine the health of the kidneys. Risk factors for kidney dysfunction such as hypertension, diabetes, cardiovascular disease, obesity, elevated cholesterol or a family history of kidney disease. It may also be when has signs and symptoms of kidney disease, though in early stage often no noticeable symptoms are observed. Kidney panel is useful for general health screening; screening patients at risk of developing kidney disease; management of patients with known kidney disease. Estimated GFR is especially important in CKD patients CKD for monitoring, it helps to identify disease at early stage in those with risk factors for CKD (diabetes, hypertension, cardiovascular disease, and family history of kidney disease). Early recognition and intervention are important in slowing the progression of CKD and preventing its complications.



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C/o : Apollo Clinic **Bill Date**

: 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM

Result Date : 20-Apr-2024 01:11 PM Report Status

: Final

Test Name

Rh Type

Result

Unit

UHID

Reference Value

: 2004240021

2004240021

Method

Blood Group & Rh Typing-Whole Blood EDTA

Blood Group

Positive

Slide/Tube

agglutination

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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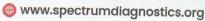
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Reg. No. : 2004240021

C/o : Apollo Clinic **Bill Date** : 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM **Result Date** : 20-Apr-2024 01:55 PM

Report Status : Final

Test Name	Result	Unit	Reference Value	Method
Complete Haemogram-Whole F	Blood EDTA			•
Haemoglobin (HB)	15.90	g/dL	Male: 14.0-17.0 Female:12.0-15.0 Newborn:16.50 - 19.50	Spectrophotmeter
Red Blood Cell (RBC)	4.94	million/cur	mm3.50 - 5.50	Volumetric Impedance
Packed Cell Volume (PCV)	45.70	%	Male: 42.0-51.0 Female: 36.0-45.0	Electronic Pulse
Mean corpuscular volume (MCV)	92.60	fL	78.0- 94.0	Calculated
Mean corpuscular hemoglobin MCH)	32.20	pg	27.50-32.20	Calculated
Mean corpuscular hemoglobin concentration (MCHC)	34.80	%	33.00-35.50	Calculated
Red Blood Cell Distribution Width SD (RDW-SD)	50.40	fL	40.0-55.0	Volumetric
Red Blood Cell Distribution CV (RDW-CV)	16.00	%	Male: 11.80-14.50 Female:12.20-16.10	Impedance Volumetric
Mean Platelet Volume (MPV)	9.90	fL	8.0-15.0	Impedance Volumetric
Platelet	2.98	lakh/cumm	1.50-4.50	Impedance Volumetric
latelet Distribution Width PDW)	10.40	%	8.30 - 56.60	Impedance Volumetric
White Blood cell Count (WBC)	6670.00	cells/cumm	Male: 4000-11000 Female 4000-11000 Children: 6000-17500 Infants: 9000-30000	Impedance Volumetric Impedance
	62.90	%	40.0-75.0	Light
ymphocytes	30.00	%	20.0-40.0	scattering/Manual Light
osinophils	4.10	%	0.0-8.0	scattering/Manual Light scattering/Manual

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Age / Gender : 41 years / Male

Ref. By Dr. : Dr. APOLO CLINIC

Reg. No. : 2004240021 C/o : Apollo Clinic **Bill Date** : 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM **Result Date** : 20-Apr-2024 01:55 PM

Report Status : Final

Test Name	Result	Unit	Reference Value	Method
Monocytes	2.70	%	0.0-10.0	Light
Basophils	0.30	%	0.0-1.0	scattering/Manual Light
Absolute Neutrophil Count	4.19	10^3/uL	2.0-7.0	scattering/Manual Calculated
Absolute Lymphocyte Count	2.00	10^3/uL	1.0-3.0	Calculated
Absolute Monocyte Count	0.18	10^3/uL	0.20-1.00	Calculated
Absolute Eosinophil Count	280.00	cells/cumm	40-440	S WAR AND THE REST OF THE PARTY
Absolute Basophil Count	0.02	10^3/uL	0.0-0.10	Calculated
Erythrocyte Sedimentation Rate (ESR)	04	mm/hr	Female: 0.0-20.0 Male: 0.0-10.0	Calculated Westergren

2004240021

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Peripheral Smear Examination-Whole Blood EDTA

Method: (Microscopy-Manual)

RBC'S : Normocytic Normochromic.

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

: Adequate in number and normal in morphology. **Platelets**

No abnormal cells or hemoparasites are present.

Impression: Normocytic Normochromic Blood picture.



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Bill Date : 20-Apr-2024 08:51 AM

Sample Col. Date: 20-Apr-2024 08:51 AM **Result Date** : 20-Apr-2024 03:15 PM

Report Status : Final

Test Name Result Unit Reference Value Method Post Prandial Urine Sugar Negative Negative Dipstick/Benedicts(Man

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