

CERTIFICATE OF MEDICAL FITNESS

NAME: Mrs. Shwefa.	
AGE/GENDER: 2948 F	
HEIGHT: 147cm	WEIGHT: 55 Kgs
IDENTIFICATION MARK:	
BLOOD PRESSURE: 120 80 100 119	
PULSE: 120 80 mily	
RS:P Normal	
ANY OTHER DISEASE DIAGNOSED IN THE PAST:	U.
ALLERGIES, IF ANY:	
LIST OF PRESCRIBED MEDICINES:	
ANY OTHER REMARKS:	
of Ms who has signed in m disease and is fit for employment.	
x Ewotha D.G. Signature of candidate	Dr. BINDURAJ. R Signature of Medical Officer
Place: Spenton Diagnostics & healt. Date: 01/4/24	b cace
Disclaimer: The patient has not been checked for COVID. covid status of the patient examined	This certificate does not relate to the
, parising	





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

DATE: 09.11.24.

EYE EXAMINATION

NAME: (Wys Jane Cha	AGE: 29	W GENDER: F/M
	RIGHT EYE	LEFT EYE
Vision	641.m	66100
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

Dr. ASHOK SARODHE Sc., M.B.B.S., D.O.M.S. Eye Consultant & Surgeon Consultant (Opthalmologist)







NAME	AGE	GENDER
MRS SHWETA	29 yrs	Sendo.

DENTAL EXAMINATION REPORT:

8	7	6	5	4	3	2	1	1	2	3	4	5	6	7	8
8	7	6	5	4	3	2	1	1	2	3	4	5	6	7	8

C: CAVITY

M: MISSING

O: OTHERS

ADVISED:

CLEANING / SCALING / ROOTS PLANNING / FLOSSING & POLISHING / OTHERS

REMARKS:

SIGNATURE OF THE DENTAL SURGEON

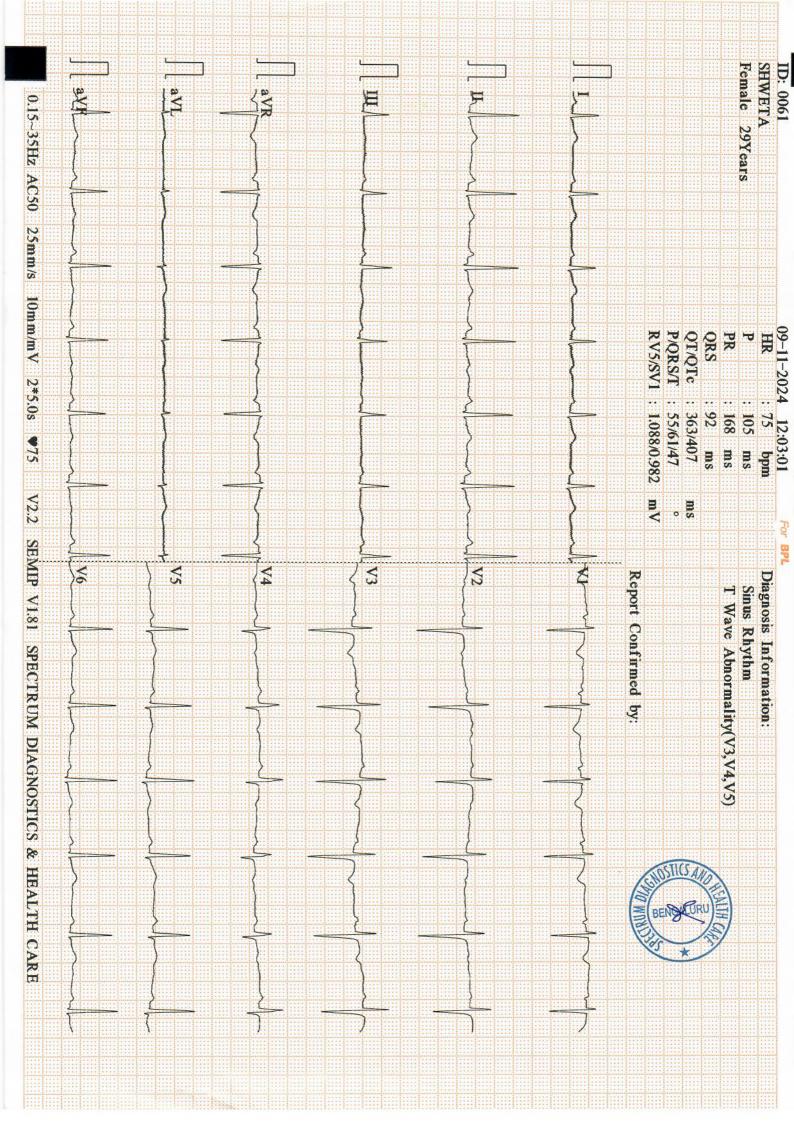
SEAL

DATE

Dr. SACHDEV NAGARKAR B.D.S., F.A.G.E., F.P.F.A. (USA) Reg. No: 2247/A









SPECTRUM DIAGNOSTCIS

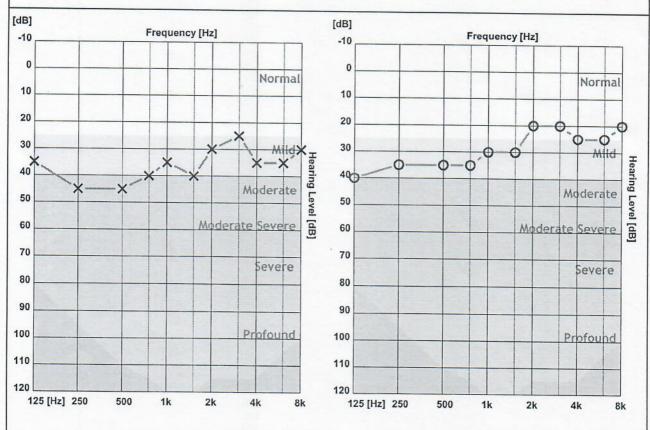
Bangalore

Patient ID: 0047 Name: SHWETHA

CR Number : 20241109104912 Registration Date : 09-Nov-2024 Age: 29

Gender: Female

Operator: spectrum diagnostics



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

	Average	High	Mid	Low
AIR Left	35.91 dB	31.25 dB	35.00 dB	41.25 dB
AIR Right	28.64 dB	22.50 dB	26.67 dB	36.25 dB

Clinical Notes:

RIGHT EAR=NORMAL LEFT EAR=NORMAL







Name

: MRS. SHWETA

: APOLLO CLINIC

Age / Gender Ref. By Dr.

: 29 years / Female : C/O APOLO CLINIC

Reg. No. C/o

: 0911240061

: 0911240061

Bill Date

: 09-Nov-2024 09:41 AM

Sample Col. Date: 09-Nov-2024 09:41 AM

Result Date

: 09-Nov-2024 12:51 PM

Report Status : Final

Test Name

Result

Unit

Reference Value

Method

CHEST PA VIEW

- · Visualised lungs are clear.
- Bilateral hila appears normal.
- · Cardia is normal in size.
- · No pleural effusion.

IMPRESSION: No significant abnormality.



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: 09 Nov, 2024 04:31 pm

DR PRAVEEN B, MBBS, DMRD, DNB Consultant Radiologist

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

+91 77604 97644 | 080 2337 1555

info@spectrumdiagnostics.org





Age / Gender : 29 years / Female Ref. By Dr. : C/O APOLO CLINIC

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C/o : APOLLO CLINIC

Bill Date

: 09-Nov-2024 09:41 AM

Sample Col. Date: 09-Nov-2024 09:41 AM

Result Date

: 09-Nov-2024 01:28 PM

Report Status : Final

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Method

2D ECHO

2D ECHO CARDIOGRAHIC STUDY M-MODE

Cardiograhic Study		Size
Aorta	28	mm
Left Atrium	26	mm
Right Ventricle	25	mm
Left ventricle (Diastole)	47	mm
Left ventricle(Systole)	20	mm
Ventricular Septum (Diastole)	07	mm
Ventricular septum (Systole)	10	mm
Posterior Wall (Diastole)	08	mm
Posterior Wall (Systole)	12	mm
Fractional Shortening	30	%
Ejection fraction	60	%

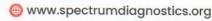
DOPPLER /COLOUR FLOW

fitral Valve Velocity MVE- 0.89m/s		MVA - 0.7	72m/s	E/A-1.23	
Tissue Doppler	e' (Septal) 10cm/s	E/e'(Septal) -8			
Velocity/ Gradient acro valve	ss the Pulmonic	0.83m/s	3mn	nHg	
Max. Velocity / Gradie valve	1.34m/s	7mn	nHg		
Velocity / Gradient acro	e 1.87 m/s	19m	mHg		

Page 1 of 2 Tejas Arcade, #9/1, 1st Main Road, Dr. Rajkumar Road, Rajajinagar, Opp. St. Theresa Hospital, Bengaluru - 560010









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Method

2DECHO Cardiographic Study

- SITUS SOLITUS, LEVOCARDIA
- SYSTEMIC VEINS: Normal drainage. IVC-1.2>50% collapse with inspiration.
- · PULMONARY VEINS: Normal drainage.
- RIGHT ATRIUM: Normal size, LEFT ATRIUM: Normal size.
- RIGHT VENTRICLE: Normal size & Adequate function.
- LEFT VENTRICLE: Normal size; No RWMA; LV Systolic function adequate.
- IAS: INTACT: IVS: INTACT.
- MITRAL VALVE: No stenosis; No regurgitation
- TRICUSPID VALVE: No stenosis; No regurgitation
- AORTIC VALVE: No stenosis; No regurgitation
- PULMONIC VALVE: No stenosis; No regurgitation
- GREAT ARTERIES: Normally related.
- · AORTA: Left aortic arch. No aortic dissection
- PULMONARY ARTERY: Confluent branch pulmonary arteries
- · NO PDA.
- · No pericardial effusion.

IMPRESSION:

- ADEQUATE LEFT VENTRICLE SYSTOLIC FUNCTION
- NO REGIONAL WALL MOTION ABNORMALITY
- ADEQUATE RIGHT VENTRICLE SYSTOLIC FUNCTION
- NO PAH



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: Durga

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: 09 Nov, 2024 01:29 pm

Ms.Durga V., ECHO Technician

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info@spectrumdiagnostics.org





NAME AND LAB NO	MRS SHWETA	REG -0061
AGE & SEX	29 YRS	FEMALE
DATE AND AREA OF INTEREST	09.11.2024	
REF BY	C/O APOLO CLINIC	

USG ABDOMEN AND PELVIS

LIVER:

Normal in size and echogenicity

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

GALL BLADDER:

Partially distended . No obvious calculus in the visualised luminal portion.

SPLEEN:

Normal in size and echotexture. No 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.

LEFT KIDNEY:

Left kidney is normal in size & echotexture

No evidence of calculus/ hydronephrosis.

URINARY BLADDER:

Well distended. No wall thickening/calculi.

UTERUS

Anteverted, Normal in size 6.9 X2.9 X3.5 cm and echotexture.

No obvious mass lesion

Endometrium is normal.ET - 6 mm.

OVARIES

Bilateral mild polycystic ovarian appearance

RO-3.2 X1.8 X2.6 cm vol 8 cc $\,$, $\,$ LO $\,$ -3.6 X 2.0 X2.6 cm vol 10 cc $\,$

No obvious adnexal mass lesions.

No evidence of ascites.

IMPRESSION:

> Bilateral mild polycystic ovarian appearance.

Suggested clinical / lab correlation

DR PRAVEEN B, DMRD, DNB CONSULTANT RADIOLOGIST







Name Age / Gender

: MRS. SHWETA

Ref. By Dr.

: 29 years / Female : C/O APOLO CLINIC

Reg. No.

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C/o

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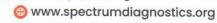
: 09-Nov-2024 12:06 PM : Final

Test Name	Result	Unit	Reference Value	Method
Complete Haemogram-Whole I	Slood EDTA			Method
Haemoglobin (HB)	14.30	-/JT	_	
Red Blood Cell (RBC)	5.31	g/dL million/cur	Female: 12.0 - 15.0 nm3.50 - 5.50	Spectrophotmeter Volumetric
Packed Cell Volume (PCV)	41.70	%	T- 1-262	Impedance
Mean corpuscular volume (MCV)	78.50	fL	Female: 36.0 - 45.0 78.0- 94.0	Electronic Pulse Calculated
Mean corpuscular hemoglobin (MCH)		pg	27.50-32.20	Calculated
Mean corpuscular hemoglobin concentration (MCHC)	34.30	%	33.00-35.50	Calculated
Red Blood Cell Distribution Width SD (RDW-SD)	33.60	fL	40.0-55.0	Volumetric
Red Blood Cell Distribution CV (RDW-CV)	14.60	%	Female: 12.20 - 16.10	Impedance Volumetric
Mean Platelet Volume (MPV)	10.00	fL	8.0-15.0	Impedance Volumetric
Platelet	4.41	lakh/cumm	1.50-4.50	Impedance Volumetric
Platelet Distribution Width PDW)	10.90	%	8.30 - 56.60	Impedance Volumetric
White Blood cell Count (WBC)	7960	cells/cumm	Female: 4000.0 - 11000.0	Impedance Volumetric
Neutrophils	59.80	%	40.0-75.0	Impedance Light
ymphocytes	33.80	%	20.0-45.0	scattering/Manual Light
osinophils	2.50	%	0.0-8.0	scattering/Manual Light
Ionocytes	3.90	%	0.0-10.0	scattering/Manual Light
asophils	0.00	%	0.0-1.0	scattering/Manual Light
bsolute Neutrophil Count	4.76	10^3/uL	2.0- 7.0	scattering/Manual Calculated













Age / Gender : 29 years / Female Ref. By Dr. : C/O APOLO CLINIC

Reg. No. : 0911240061

C/o: APOLLO CLINIC **Bill Date**

Female: 0.0 - 20.0

: 09-Nov-2024 09:41 AM

Westergren

Sample Col. Date: 09-Nov-2024 09:41 AM

Result Date : 09-Nov-2024 12:06 PM Report Status : Final

Test Name	_			
1 est ivaine	Result	Unit	Reference Value	Method
Absolute Lymphocyte Count	2.69	10^3/uL	1020	
Absolute Monocyte Count	0.31	10^3/uL	1.0-3.0	Calculated
Absolute Eosinophil Count	200.00	cells/cumm	0.20-1.00	Calculated
Absolute Basophil Count	0.00	10^3/uL	40-440	Calculated
Erythrocyte Sedimentation	19	mm/hr	0.0-0.10	Calculated

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mm/hr

Peripheral Smear Examination-Whole Blood EDTA

Method : (Microscopy-Manual)

Rate (ESR)

RBC'S : Normocytic Normochromic.

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

: Adequate in number and normal in morphology. No abnormal cells or hemoparasites are present.

Impression: Normocytic Normochromic Blood Picture.



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Name Age / Gender

: MRS. SHWETA

Ref. By Dr.

: 29 years / Female : C/O APOLO CLINIC

Reg. No. C/o

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Bill Date

: 09-Nov-2024 09:41 AM Sample Col. Date: 09-Nov-2024 09:41 AM

Result Date

: 09-Nov-2024 12:42 PM

Report Status : Final

Test Name	Result	Unit	Reference Value	Method
LFT-Liver Function Test -Serun	<u>m</u>		,	ALCHOU
Bilirubin Total-Serum	1.18	mg/dL	0.2-1.0	Caffeine
Bilirubin Direct-Serum	0.29	mg/dL	0.0-0.2	Benzoate Diazotised Sulphanilic
Bilirubin Indirect-Serum Aspartate Aminotransferase (AST/SGOT)-Serum	0.89 24.00	mg/dL U/L	0.0-1.10 15.0-37.0	Acid Direct Measure UV with Pyridoxal - 5 -
Alanine Aminotransferase (ALT/SGPT)-Serum	33.00	U/L	Male:16.0-63.0 Female:14.0-59.0	Phosphate UV with Pyridoxal - 5 -
Alkaline Phosphatase (ALP)- Serum	109.00	U/L	Adult: 45.0-117.0 Children: 48.0-445.0	Phosphate PNPP,AMP- Buffer
Protein, Total-Serum	9.48	g/dL	Infants: 81.90-350.30 6.40-8.20	Biuret/Endpoint-
Albumin-Serum	5.01	g/dL	3.40-5.00	With Blank Bromocresol
Globulin-Serum Albumin/Globulin Ratio-Serum	4.47 1.12	g/dL Ratio	2.0-3.50 0.80-2.0	Purple Calculated Calculated



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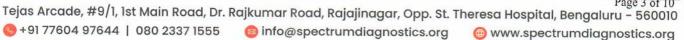
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Age / Gender : 29 years / Female Ref. By Dr. : C/O APOLO CLINIC

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Bill Date UHID : 0911240061

: 09-Nov-2024 09:41 AM

Sample Col. Date: 09-Nov-2024 09:41 AM Result Date : 09-Nov-2024 12:42 PM

Report Status : Final

Test Name	Result	Unit			
	Result		Reference Value	Method	
Fasting Urine Glucose-Urine	Negative		Negative	Dipstick/Benedicts (Manual)	
Fasting Blood Sugar (FBS)- Plasma	76	mg/dL	60.0-110.0	Hexo Kinase	

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

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

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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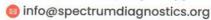
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Age / Gender Ref. By Dr.

: 29 years / Female

Reg. No.

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Test Name	Result	Unit	Reference Value	Method	
Glycosylated Haemoglobin (HbA1c)-Whole Blood EDTA			,	THE CHILD	
Glycosylated Haemoglobin (HbA1c)	5.50	%	Non diabetic adults :<5.7 At risk (Prediabetes) : 5.7 - 6.4	HPLC	
			Diagnosing Diabetes :>= 6.5 Diabetes		
			Excellent Control: 6-7		
			Fair to good Control: 7-8		
			Unsatisfactory Control :8-10 Poor Control :>10		
Estimated Average Glucose(eAG)	111.14	mg/dL		Calculated	

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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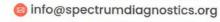
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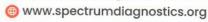
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Method

Blood Group & Rh Typing-Whole Blood EDTA

Blood Group

Rh Type

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

Gamma-Glutamyl Transferase 36.00

(GGT)-Serum

U/L

Male: 15.0-85.0

Other g-Glut-

Female: 5.0-55.0

3-carboxy-4

nitro

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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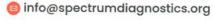
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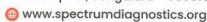
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Test Name	Result	Unit	Reference Value	Method
Thyroid function tests (TFT) Serum)-			
Tri-Iodo Thyronine (T3)-Ser	rum 1.00	ng/mL	0.60-1.81	Chemiluminescence Immunoassay
Thyroxine (T4)-Serum	11.9	μg/dL	5.50-12.10	(CLIA) Chemiluminescence
Thyroid Stimulating Hormor (TSH)-Serum	ne 1.93	μIU/mL	0.35-5.50	Immunoassay (CLIA) Chemiluminescence Immunoassay (CLIA)

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

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.

els: Graves disease, Autonomous thyroid hormone secretion, TSH defic

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Dr. Nithun Reddy C,MD,Consultant Pathologist

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Age / Gender : 29 years / Female Ref. By Dr. : C/O APOLO CLINIC

Reg. No. : 0911240061

C/o : APOLLO CLINIC Bill Date

: 09-Nov-2024 09:41 AM Sample Col. Date: 09-Nov-2024 09:41 AM

Result Date

: 09-Nov-2024 01:17 PM

Report Status : Final

Test Name	Result	Unit	Reference Value	Method
Lipid Profile-Serum		-		Action
Cholesterol Total-Serum	139.00	mg/dL	0.0-200	Cholesterol
Triglycerides-Serum	62.00	mg/dL	0.0-150	Oxidase/Peroxidase Lipase/Glycerol
High-density lipoprotein (HDL) Cholesterol-Serum	49.00	mg/dL	40.0-60.0	Dehydrogenase Accelerator/Selective
Non-HDL cholesterol-Serum Low-density lipoprotein (LDL) Cholesterol-Serum	90 78	mg/dL mg/dL	0.0130 0.0-100.0	Detergent Calculated Cholesterol esterase
Very-low-density lipoprotein (VLDL) cholesterol-Serum	12	mg/dL	0.0-40	and cholesterol oxidase Calculated
Cholesterol/HDL Ratio-Serum	2.84	Ratio	0.0-5.0	Calculated

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UHID

Interpretation:

Desirable	Poudouline III 1		_
	Borderline High	High	Very High
<200	200-239	>240	
<150	150-199	200,400	500
<130			>500
	160-189	190-219	>220
<100	100-129	160-189	>190
	Desirable <200 <150 <130 <100	<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	
Kidney Function Test (KFT)-BUN,CREA,Uric Acid,Na,K,Cl-Serum Kidney Function Test (KFT)- Serum							
Blood Urea Nitrogen (BUN)	7.90	mg/dL	7.0-18.0			GLDH,Kinetic	
Creatinine-Serum	0.56	mg/dL	Male: 0.70-1.30 Female: 0.55-1.02			Assay Modified	
Uric Acid-Serum Electrolytes	5.11	mg/dL	Male: 3.50-7.20 Female: 2.60-6.0	ts 18		kinetic Jaffe	
Sodium (Na+)-Serum Potassium (K+)-Serum Chloride (Cl-)-Serum	136.9 4.69 101.30	mmol/L mmol/L mmol/L	135.0-145.0 3.50-5.50 96.0-108.0	**		ISE-Direct ISE-Direct ISE-Direct	

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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Name

: MRS. SHWETA

Age / Gender Ref. By Dr.

: 29 years / Female : C/O APOLO CLINIC

Reg. No.

: 0911240061

C/o

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Test Name

Result

Unit

Reference Value

Method

Postprandial Urine glucose-Urine

Negative

Negative

Dipstick/Benedicts

(Manual)

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

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

Calcium, Total- Serum

10.00

mg/dL

8.50-10.10

Spectrophotometry

(O-

Cresolphthalein complexone)



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