



Veena Nagar, Phase 2, Tulsi Pipeline Road, Mulund (West), Mumbai - 400080. email: medical.admin_ahm@apexhospitals.in | www.apexgroupofhospitals.com

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Land Line No. 022 - 4162 4000 (100 Lines) Reception No.: 8422854005

DEPARTMENT OF LABORATORY SCIENCES

Patient Name Mrs. VANITA SHAMBHAG LabNo **UHID/IP No** 120065755 / 526 Sample Date 18/05/2024 3:05PM Age/Gender 59 Yrs/Female **Receiving Date** 18/05/2024 3:15PM Bed No/Ward 18/05/2024 6:30PM OPD Report Date Prescribed By Final Dr. Apex Hospitals **Report Status**

HAEMATOLOGY

Test Name	Result	Unit	Biological Ref. Range	Method
COMPLETE BLOOD COUNT(CBC) Sample: W. B. EDTA	EDTA WHOLE	BLOOD		
Haemoglobin Estimation (Hb)	12.3 L	gm/dl	12.5 - 16.0	SLS- Hb Method
RBC Count (Red Blood Cell)	4.56	10^6/uL	4.50 - 6.50	
PCV (Haematocrit)	37.8	%	36.0 - 46.0	
MCV	82.89	fl	78 - 95	Calculated
MCH	26.97	pg	26 - 31	Calculated
MCHC	32.54	gm/dl	30 - 36	Calculated
RDW	15.2	%	11.0 - 16.0	Calculated
Total Leukocyte Count (TLC)	7000	cells/cu.mr	n 4000.0 - 11000.0	
Neutrophil %	55	%	40 - 75	
Lymphocyte %	40	%	20 - 45	
Eosinophil %	02	%	0 - 6	
Monocytes %	03	%	1 - 10	
Basophil %	00	%	0 - 2	
WBCs Morphology	No Abnorma	lity Detected		
RBCs Morphology	Hypochromia	a		
Platelet Count	225	10^3/uL	150 - 450	DC Detection
Platelets Morphology	Adequate on	smear		
MPV	9.3	fl	7 - 12	
ERYTHROCYTE SEDIMENTATION Sample: W. B. EDTA	N RATE (ESR)			
ESR (Erythrocyte Sed.Rate)	18	mm/hr	< 20	Westergren

-- End Of Report--

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APEX HOSPITALS MULUND



Veena Nagar Phase II, Tulsi Pipe Line Road, Near Swapna Nagri Road, Mulund (W) Mumbai 400 080. email: info@apexhospitals.in | www.apexgroupofhospitals.com



Tele .: 022-41624000 (100 Lines)

DEPARTMENT OF LABORATORY SCIENCES

Patient Name Mrs. VANITA SHAMBHAG **UHID/IP No** 120065755 / 526 Age/Gender 59 Yrs/Female

Bed No/Ward OPD

Prescribed By

Dr. Apex Hospitals

LabNo

Sample Date

18/05/2024 3:05PM

Receiving Date Report Date

18/05/2024 3:15PM

18/05/2024 6:30PM

Report Status

Final

IMMUNO-HAEMATOLOGY

Test Name	Result	Unit	Biological	Method
			Ref. Range	

BLOOD GROUPING

Sample: W. B. EDTA

Blood Group (ABO and Rh)

"B" Rh Positive

-- End Of Report--







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Reception No.: 8422854005

DEPARTMENT OF LABORATORY SCIENCES

	Mrs. VANITA SHA			LabNo	1582	
i	120065755 / 526	5		Sample Date	18/05/2024	
• .	59 Yrs/Female OPD			Receiving Date	18/05/2024 18/05/2024	
· '	ספט Dr. Apex Hospita	de		Report Date Report Status	Final	0.30014
Prescribed by	DI. Apex Hospita			Keport Status		
LDL Cholesterol : HD Ratio	DL Cholesterol	2.77		0.00 - 4.50		Calculated Value
LIVER FUNCTION T Sample: Serum	EST (LFT) SERU	JM				
Bilirubin Total (TBil)		0.78	mg/dl	0.30 - 1.30		Diphyline Diazonium Salt
Bilirubin Direct (Dbil)	0.22	mg/dl	0.00 - 0.50		
Bilirubin indirect		0.56	mg/dl	0 - 1		
SGPT (ALT)1		14.2	U/L			IFCC modified
SGOT (AST)		25.2	U/L			IFCC modified
Protein Total		6.4	gm/dl	6.00 - 8.00		Biuret
Albumin		3.8	gm/dl	3.40 . 4.90		
Globulin		2.60	gm/dl	2.30 - 3.60		Calculated Value
A/G Ratio (Albumin/	Globulin Ratio)	1.46		1.00 - 2.50		Calculated Value
Alkaline Phosphatase	е	143.1	IU/L	64 - 306		
RFT (RENAL FUNCT: Sample: Serum	ION TEST)					
Creatinine		0.81	mg/dl	0.60 - 1.40		
UREA		26.1	mg/dl	10 - 50		CDC Urease,Colorimetric
BUN - Blood Urea Ni	trogen	12.2	mg/dl			
Calcium		8.9				
Uric Acid		6.43 H	mm/h	r 2.6 - 6.0		URICASE- PEROXIDASE
Phosphorus		3.8	mg/dl	2.5 - 4.5		Phosphomolybdate Reduction
Sodium		141.6	mmol/	L 135 - 146		ISE Direct
Potassium		4.64	mmol/	L 3.5 - 5.5		ISE Direct
Chloride		110.4 H	mmol/	'L 95 - 109		ISE Direct
Protein Total		6.4	gm/dl	6.00 - 8.00		Biuret
Albumin		3.8	gm/dl	3.40 . 4.90		
Globulin		2.60	gm/dl	2.30 - 3.60		Calculated Value
A/G Ratio (Albumin/	Globulin Ratio)	1.46				Calculated Value

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BIOCHEMISTRY

Test Name	Result	Unit	Biological Ref. Range	Method	
GLUCOSE (PP) Sample: Fl. Plasma					
Blood Sugar(2 Hours PP)	95.2				
Urine PP Sugar	SNR				
Urine PP Ketone	SNR				

Note: An individual may show higher fasting glucose level in comparison to post prandial glucose level due to

following reasons :

The glycaemic index and response to food consumed, Changes in body composition, Increased insulin response and

sensitivity, Alimentary hypoglycemia, Renal glycosuria, Effect of oral hypoglycaemics & Insulin treatment.

GLUCOSE (FASTING)

Sample:	FI. Plasn	าล			
Chicoco	/Cashina	Dlood	Cuank	/ EBC)	

Glucose (Fasting Blood Sugar / FBS) 78.1 Urine Fasting Sugar SNR Urine Fasting Ketone SNR

LIPID PROFILE SERUM

Sample: Serum				
Cholesterol-Total	185.1	mg/dl	< 200.00	Cholesterol Oxidase,Esterase,Pero xidase
Triglycerides	95.2	mg/dl	< 150	Enzymatic End point
HDL Cholesterol	44.1	mg/dl	30.00 - 70.00	Phosphotungstat
VLDL Cholesterol	19.04	mg/dl	6.00 - 35.00	Calculated Value
LDL Cholesterol	121.96	mg/dl	< 160.00	Calculated Value
Cholesterol Total : HDL Cholesterol Ratio	4.20		0.00 - 4.80	Calculated Value

C2____





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Reported On

Land Line No. 022 - 4162 4000 (100 Lines) Reception No.: 8422854005

Patient ID : 2405075040

Patient Name

: MRS. VANITA SHAMBHAG

Age

: 59 Yrs

Gender

טיין אינים

Ref. By Doctor

: FEMALE : APEX HOSPITAL

Sample Collected At: APEX HOSPITAL MULUND

For Authenticity Scan QR Code

Registered On : 18/05/2024,05:03 PM Collected On : 19/05/2024,06:18 AM

: 19/05/2024,08:27 AM

Glycosylated Hemoglobin (GHb/HBA1c)

Test Name	Result	Unit	Biological Reference Interval
HbA1c (Glycocylated Haemoglobin)	6.50	%	Below 6.0% : Normal
			6.0% 7.0% : Good Control
			7.0% - 8.0% : Fair Control
			8.0%-10% : Unisatisfactory
			Above 10% Poor Control
HPLC- H9			
Mean Blood Glucose Calculated	139.8	mg/dL	70 - 125

CLINICAL SIGNIFICANCE:

Glycosylated Haemoglobin is a acurate and true index of the "Mean Blood Glucose Level" in the body for the previous 2-3 months.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.

Factors affecting HbA1c results:

Increased in: High fetal hemoglobin, Chronic renal failure, Iron deficiency anemia, Splenectomy, Increased serum triglycerides, Alcohol ingestion, Lead/opiate poisoning and Salicylate treatment.

Decreased in: Shortened RBC lifespan (Hemolytic anemia, blood loss), following transfusions, pregnancy, ingestion of large amount of Vitamin E or Vitamin C and Hemoglobinopathies

Reflex tests: Blood glucose levels, CGM (Continuous Glucose monitoring)

----- End of Report --

Results relate only to the sample as received. Kindly correlate with clinical condition

Note: If the test results are alarming or unexpected, Client is advised to contact the Physician immediately for possible remedial action.

This report is system generated and electronically authenticated.

Page 1 of 1

Dr. Roshan Shaikh MBBS MD Pathology Consultant Pathologist





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Reception No.: 8422854005

Mrs. VANITA SHAMBHAG

DOB

Age

CRM

Gender Female

59 Years

18-05-2024 17:00 Collected :

18-05-2024 22:38 Received 18-05-2024 23:37 Reported

Status Final

Sample Quality

Lab ID

Adequate Location

MUMBAI Ref Bv APEX HOSPITAL

SANIAY PANDEY -MU058 Client

Unit Parameter Biological Ref. Interval Result

THYROID FUNCTION TEST

Tri Iodo Thyronine (T3 Total), Serum CLIA

1.05

ng/mL

0.4 - 1.81

Clinical significance:-

Triiodothyronine (T3) values above 3.07 ng/mL in adults or over age related cutoffs in children are consistent with hyperthyroidism or increased thyroid hormone-binding proteins. Abnormal levels (high or low) of thyroid hormone-binding proteins (primarily albumin and thyroid-binding globulin) may cause abnormal T3 concentrations in euthyroid patients. Please note that Triiodothyronine (T3) is not a reliable marker for hypothyroidism. Therapy with amiodarone can lead to depressed T3 values.

Thyroxine (T4), Serum

9.92

µg/dL

5.5-11.0

Clinical significance:-

Thyroxine (T4) is synthesized in the thyroid gland. High T4 are seen in hyperthyroidism and in patients with acute thyroiditis. Low T4 are seen in hypothyroidism, myxedema, cretinism, chronic thyroiditis, and occasionally, subacute thyroiditis. Increased total thyroxine (T4) is seen in pregnancy and patients who are on estrogen medication. These patients have increased total T4 levels due to increased thyroxine-binding globulin (TBG) levels. Decreased total T4 is seen in patients on treatment with anabolic steroids or nephrosis (decreased TBG levels).

Thyroid Stimulating Hormone (TSH), Serum

CLIA

2.125

ulU/mL

Nonpregnant: 0.4 - 5.5

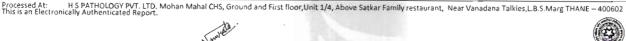
Pregnancy:

First Trimester: 0.3-4.5 Second Trimester: 0.5-4.6 Third trimester: 0.8-5.2

Clinical significance:

In primary hypothyroidism, TSH (thyroid-stimulating hormone) levels will be elevated. In primary hypothyroidism, TSH levels will be low. TSH estimation is especially useful in the differential diagnosis of primary (thyroid) from secondary (pituitary) and tertiary (hypothalamus) hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low or normal. Elevated or low TSH in the context of normal free thyroxine is often referred to as subclinical hypo- or hyperthyroidism, respectively

Pregnancy	American Thyroid	American European	Thyroid society
	Association	Endocrine	Association
1st trimester	< 2.5	< 2.5	< 2.5
2nd trimester	< 3.0	< 3.0	< 3.0
3rd trimester	< 3.5	< 3.0	< 3.0







Age

Gender

CRM

59 Years

Female

APEX HOSPITALS MULUND DIAGNOSTIC





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Lab ID Mrs. VANITA SHAMBHAG Collected 18-05-2024 17:00 DOB 18-05-2024 22:38 Sample Quality Received

Status

18-05-2024 23:37 Reported

Location

Adequate MUMBAI

Final

APEX HOSPITAL Ref Bv

Client

SANJAY PANDEY -MU058

Parameter	Result	Unit	Biological Ref. Interval	
		/ 1	420.044	
Vitamin B12, Serum	179.00	pg/mL	120-914	

Clinical significance:

Vitamin B12 (cobalamin) is necessary for hematopoiesis and normal neuronal function. The body uses its vitamin B12 stores very economically, reabsorbing vitamin B12 from the ileum and returning it to the liver; very little is excreted. Vitamin B12 deficiency may be due to lack of IF secretion by gastric mucosa (eg. gastrectomy, gastric atrophy) or intestinal malabsorption (eg, ileal resection, small intestinal diseases). Pernicious anemia is a macrocytic anemia caused by vitamin B12 deficiency that is due to a lack of IF secretion by gastric mucosa. Serum methylmalonic acid and homocysteine levels are also elevated in vitamin B12 deficiency states.





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Mrs. VANITA SHAMBHAG

DOB

Age

CRM

Gender

59 Years

Female

Collected 18-05-2024 17:00

18-05-2024 22:38 Received 18-05-2024 23:40 Reported

Status Final Lab ID

Sample Quality

MUMBAI

Adequate

>80: Toxicity possible

Location Ref By APEX HOSPITAL

Client SANJAY PANDEY -MU058

Parameter	Result	Unit	Biological Ref. Interval
Vitamin D - 25-Hydroxy, Serum CLIA	27.48	ng/mL	<10: Severe deficiency 10-19: Mild to moderate deficiency 20-50: Optimum level 51-80: Increased risk of hypercalciuria

Clinical significance:-

A low blood level of 25-hydroxyvitamin D may mean that a person is not getting enough exposure to sunlight or enough dietary vitamin D to meet his or her body's demand or that there is a problem with its absorption from the intestines. Occasionally, drugs used to treat seizures, particularly phenytoin (Dilantin), can interfere with the production of 25-hydroxyvitamin D in the liver. There is some evidence that vitamin D deficiency may increase the risk of some cancers, immune diseases, and cardiovascular disease. A high level of 25-hydroxyvitamin D usually reflects excess supplementation from vitamin pills or other nutritional supplements.

End Of Report -



