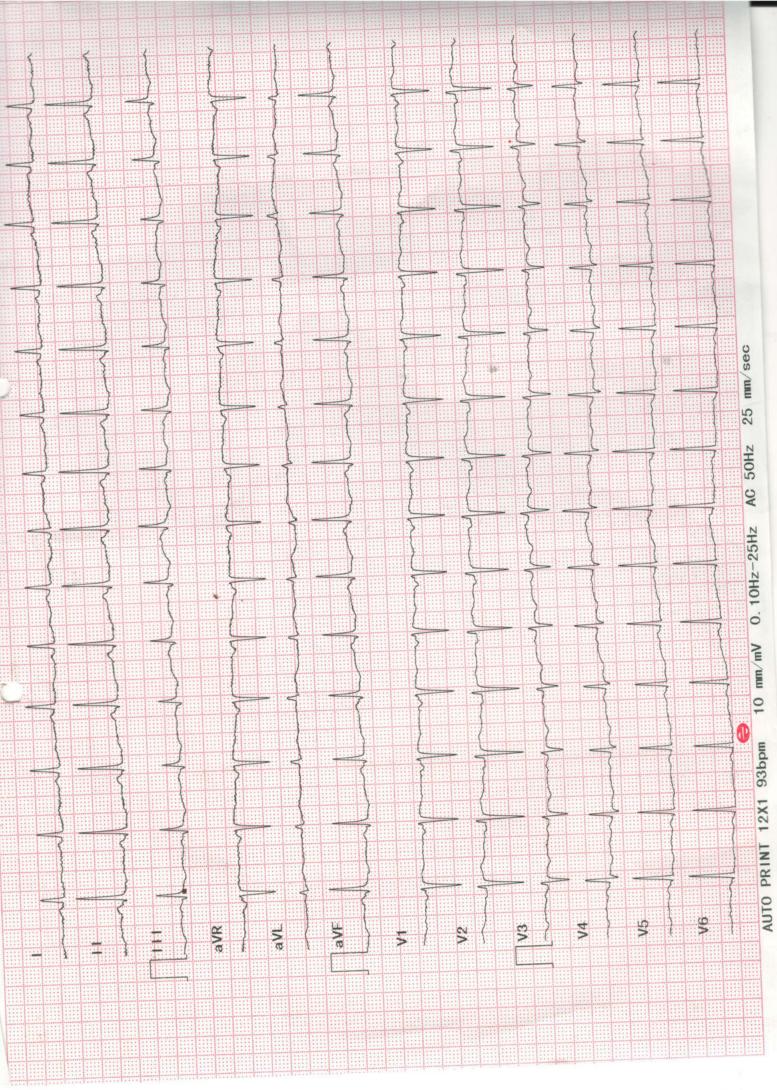
Dash	
Jayshree	Var 14 0 215
Dr.	b: www.uni-em.com. TMT Var
	em@electromedicals.net; We
	Fax: +91-731-4031180, E-Mail:
	1 "CENNEND-TE/-TE/ :
Tom output	.Tat .atonin

UNI-EM, Ind Technician : VIKESH JADHAV

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MUMBAI	TEST REPORT		••	••	••	6	• 1 • 1	Arriver		< <	120 / 8	<b>``</b>	、 \	<b>、</b>	~	1			ACON VAN	rate 187 hhm										
MUMBAI	TREADMILL	PROTOCOL	HISTORY	INDICATION	MEDICATION	Q 11	п.к. bom		Ͽ	104	130	101	ILI	118	135	122	-			heart										
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						CDFFD	Km/Hr				с с	4.1	5.4						ä	bpm 91	/ 100	ACHIEVED					IONOTROPIC RES			SUBRCC TECH TC MECANTINE TO TIMITAL TANK
		N					TIME				0:35 ว.ธธ	2:55	0:18	2:20	2:55	5:55 2:55	0:33		: 6:15	: 171	••	••	••							- aca attu
	MEGHA TANK	12-11-202	33 /F	136 / /l Salf	4422	TOTAT.	TIME				2.55	5:55	6:18	8:53	9:28	12:28	0 1 1		DURATION	MAX HEART RATE	PRESSURE	TERMINATION	SE	ONSP	LONS	GOOD EFFORT TOLERANCE	NORMAL CHRONOTROPIC AND	NO ANGINA / ARRHYTHMIA	CHANGES.	KUAN OL UD
	MCS. MEGH	DATE	AGE/SEX	REF. BY		PHASE												RESULTS	EXERCISE DURATION	MAX HEARD	MAX BLOOD	REASON OF	BP RESPONSE	ARRYTHMIA H.R. RESPONSE	IMPRESSIONS	GOOD REFO	NORMAL CH	NO ANGINA	NO ST - T CHANGES	ampred mr
						2H4			RECOVERY	STANDING	Stage 1	age 2	<b>PK-EXERCISE</b>	COVERY	KECOVERY	KECOVERY RECOVERY														

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## **DIAGNOSTICS REPORT**

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Order Date	: 12/11/2022 10:29
Age/Sex	: 33 Year(s)/Female	Report Date	: 12/11/2022 12:35
UHID	: SHHM.52556	IP No	:
Ref. Doctor	: Self	Facility	: SEVENHILLS HOSPITAL, MUMBAI

## 2D ECHOCARDIOGRAPHY WITH COLOUR DOPPLER STUDY

Normal LV and RV systolic function.

Estimated LVEF = 60%

No LV regional wall motion abnormality at rest .

All valves are structurally and functionally normal.

Normal sized cardiac chambers.

No LV Diastolic dysfunction .

No pulmonary arterial hypertension.

No regurgitation across any other valves.

Normal forward flow velocities across all the cardiac valves.

Aorta and pulmonary artery dimensions: normal.

IAS / IVS: Intact.

No evidence of clot, vegetation, calcification, pericardial effusion. COLOUR DOPPLER: NO MR/AR.



Dr.Jayashree Dash,

(Junior Consultant NIC)

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

## **Blood Bank**

Test Name		Resu	ılt					
Sample No :	O0248609A	Collection Date :	12/11/22 1	0:33 Acl	k Date :	12/11/2022 10:52	Report Date :	12/11/22 11:37
BLOOD GR	OUPING (ABO+F	RH) BY COLUMN	AGGLUT	INATION N	METHOD	þ		
BLOOD GRO	UP (ABO)	'0'						
Rh Type		POSI	TIVE					
results per	<u>The reported</u> tain to the samp the blood centre							

Interpretation :

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Blood typing is used to determine an individual's blood group, to establish whether a person is blood group A, B, AB, or O and whether he or she is Rh positive or Rh negative. Blood typing has the following significance,

• Ensure compatibility between the blood type of a person who requires a transfusion of blood or blood components and the ABO and Rh type of the unit of blood that will be transfused.

• Determine compatibility between a pregnant woman and her developing baby (fetus). Rh typing is especially important during

pregnancy because a mother and her fetus could be incompatible.

• Determine the blood group of potential blood donors at a collection facility.

• Determine the blood group of potential donors and recipients of organs, tissues, or bone marrow, as part of a workup for a transplant procedure.

End of Report

Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept.

[			
Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Biochemistry									
Test Name	Result	t			Unit	Ref.	Range		
Sample No: 00248609A	Collection Date :	12/11/22 10:33	Ack Date :	12/11/2022 10:50	Report I	Date :	12/11/22 11:27		
<u>GLYCOSLYATED HAEMOG</u> (HBA1C)	<u>LOBIN</u>								
HbA1c	5.61				%	6.0 contr 7.0 contr 8.0 contr	8.0% Fair to good ol 10% Unsatisfactory		
Method - BIOCHEMISTRY									
Estimated Average Glucose ( Method - Calculated	eAG) 114.31				mg/dl	90 - 3	126		
<ul> <li>Method - Calculated</li> <li>NOTES :- <ol> <li>HbA1c is used for monitoring diabetic control. It reflects the mean plasma glucose over three months</li> <li>HbA1c may be falsely low in diabetics with hemolytic disease. In these individuals a plasma fructosamine level may be used which evaluates diabetes over 15 days.</li> <li>Inappropriately low HbA1c values may be reported due to hemolysis, recent blood transfusion, acute blood loss, hypertriglyceridemia, chronic liver disease. Drugs like dapsone, ribavirin, antiretroviral drugs, trimethoprim, may also cause interference with estimation of HbA1c, causing falsely low values.</li> <li>HbA1c may be increased in patients with polycythemia or post-splenectomy.</li> <li>Inappropriately higher values of HbA1c may be caused due to iron deficiency, vitamin B12 deficiency, alcohol intake, uremia, hyperbilirubinemia and large doses of aspirin.</li> <li>Trends in HbA1c are a better indicator of diabetic control than a solitary test.</li> <li>Any sample with &gt;15% HbA1c should be suspected of having a hemoglobin variant, especially in a non-diabetic patient. Similarly, below 4% should prompt additional studies to determine the possible presence of variant hemoglobin.</li> <li>HbA1c target in pregnancy is to attain level &lt; 7.5 %.</li> <li>Method : turbidimetric and blates Associations. Standards of Medical Care in Diabetes 2015</li> </ol> </li> </ul>									
Sample No: 00248609B	Collection Date : 1	12/11/22 10:33	Ack Date :	12/11/2022 11:06	Report I	Date :	12/11/22 11:27		
GLUCOSE-PLASMA-FASTI	NG								
Glucose,Fasting	93.2				mg/dl	70 - 3	110		

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

American Diabetes Association Reference Range :

Normal : < 100 mg/dl Impaired fasting glucose(Prediabetes) : 100 - 126 mg/dl Diabetes : >= 126 mg/dl

References: 1)Pack Insert of Bio system 2) TIETZ Textbook of Clinical chemistry and Molecular Diagnostics Edited by: Carl A.burtis, Edward R. Ashwood,David e. Bruns

Interpretation :-

Conditions that can result in an elevated blood glucose level include: Acromegaly, Acute stress (response to trauma, heart attack, and stroke for instance), Chronic kidney disease, Cushing syndrome, Excessive consumption of food, Hyperthyroidism, Pancreatitis. A low level of glucose may indicate hypoglycemia, a condition characterized by a drop in blood glucose to a level where first it causes nervous system symptoms (sweating, palpitations, hunger, trembling, and anxiety), then begins to affect the brain (causing confusion, hallucinations, blurred vision, and sometimes even coma and death). A low blood glucose level (hypoglycemia) may be seen with:Adrenal insufficiency, Drinking excessive alcohol, Severe liver disease, Hypopituitarism, Hypothyroidism, Severe infections, Severe heart failure, Chronic kidney (renal) failure, Insulin overdose, Tumors that produce insulin (insulinomas), Starvation.

Sample No :	O0248609C	Collection Date :	12/11/22 10:33	Ack Date :	12/11/2022 11:01	Report Date :	12/11/22 11:27
Lipid Profile	<u>e</u>						

Total Cholesterol	207.94	mg/dl	Reference Values : Up to 200 mg/dL - Desirable 200-239 mg/dL - Borderline HIgh >240 mg/dL - High
Triglycerides	88.98	mg/dl	Reference Values: Up to 150 mg/dL - Normal 150-199 mg/dL - Borderline High 200-499 mg/dL - High >500 mg/dL - Very High
Method - Enzymatic			
HDL Cholesterol Method - Enzymatic immuno inhibition	34.68	mg/dl	0 - 60

Patient Name: Mrs. MEGHNA BH.UHID: SHHM.52556Episode: OPRef. Doctor:	AVESH TANK	Age/Sex Order Date Mobile No DOB Facility	: 12/11/20 : 9699767 : 09/06/1	177
LDL Cholesterol Method - Calculated	<b>155.46 ▲</b>		mg/dl	0 - 130
VLDL Cholesterol Method - Calculated	17.80		mg/dl	0 - 40
Total Cholesterol / HDL Cholesterol Ratio - Calculated Method - Calculated	6.00 ▲		RATIO	0 - 5
LDL / HDL Cholesterol Ratio - Calculated Method - Calculated	4.48 ▲		RATIO	0 - 4.3
References: 1)Pack Insert of Bio system 2) TIETZ Textbook of Clinical chemistry au Interpretation 1. Triglycerides: When triglycerides are ver adults. Triglycerides change dramatically in	v high greater than 1000 mg/dL, th	nere is a risk of developing pancreatiti	is in children and	

adults. Triglycerides change dramatically in response to meals, increasing as much as 5 to 10 times higher than fasting levels just a few hours after eating. Even fasting levels vary considerably day to day. Therefore, modest changes in fasting triglycerides measured on different days are not considered to be abnormal. 2. HDL-Cholesterol: HDL- C is considered to be beneficial, the so-called "good" cholesterol, because it removes excess cholesterol from

2. HDL-Cholesterol: HDL- C is considered to be beneficial, the so-called "good" cholesterol, because it removes excess cholesterol from tissues and carries it to the liver for disposal. If HDL-C is less than 40 mg/dL for men and less than 50 mg/dL for women, there is an increased risk of heart disease that is independent of other risk factors, including the LDL-C level. The NCEP guidelines suggest that an HDL cholesterol value greater than 60 mg/dL is protective and should be treated as a negative risk factor.

3. LDL-Cholesterol: Desired goals for LDL-C levels change based on individual risk factors. For young adults, less than 120 mg/dL is acceptable. Values between 120-159 mg/dL are considered Borderline high. Values greater than 160 mg/dL are considered high. Low levels of LDL cholesterol may be seen in people with an inherited lipoprotein deficiency and in people with hyperthyroidism, infection, inflammation, or cirrhosis.

#### Uric Acid (Serum)

4.7

Method - Uricase

2.6 - 6

mg/dl

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

References:

1)Pack Insert of Bio system

2) TIETZ Textbook of Clinical chemistry and Molecular DiagnosticsEdited by: Carl A.burtis, Edward R. Ashwood, David e. Bruns

Interpretation:-

Uric acid is produced by the breakdown of purines. Purines are nitrogen-containing compounds found in the cells of the body, including our DNA. Increased concentrations of uric acid can cause crystals to form in the joints, which can lead to the joint inflammation and pain characteristic of gout. Low values can be associated with some kinds of liver or kidney diseases, Fanconi syndrome, exposure to toxic compounds, and rarely as the result of an inherited metabolic defect (Wilson disease).

#### Liver Function Test ( LFT )

SGOT (Aspartate Transaminase) - SERUM Method - IFCC	25.99	U/L	0 - 31
SGPT (Alanine Transaminase) - SERUM Method - IFCC	37.96 ⊾	U/L	0 - 34
Total Bilirubin - SERUM Method - Diazo	1.25	mg/dl	0 - 2
Direct Bilirubin SERUM Method - Diazotization	0.4	mg/dl	0 - 0.4
Indirect Bilirubin - Calculated Method - Calculated	0.85 🔺	mg/dl	0.1 - 0.8
Alkaline Phosphatase - SERUM Method - IFCC AMP Buffer	79.96	U/L	0 - 105
Total Protein - SERUM Method - Biuret	7.53	gm/dl	6 - 7.8
Albumin - SERUM Method - Bromo Cresol Green(BCG)	4.45	gm/dl	3.5 - 5.2

Patient Name	: Mrs. MEGHNA B	HAVESH TANK	Age/Sex	: 33 Year	(s) / Female
UHID	: SHHM.52556		Order Date	: 12/11/2	022 10:29
Episode	: OP				
Ref. Doctor	:		Mobile No	: 969976	7177
			DOB	: 09/06/	1989
			Facility	: SEVENH	IILLS HOSPITAL, MUMBAI
Globulin - Calc Method - Calcu		3.08		gm/dl	2 - 4
A:G Ratio Method - Calcul	lated	1.44		:1	1 - 3

Gamma Glutamyl Transferase (GGT) - Gglutamyl carboxy nitroanilide - SERUM	32.43	U/L	0 - 38

Method - G glutamyl carboxy nitroanilide

#### References:

1)Pack Insert of Bio system

2) TIETZ Textbook of Clinical chemistry and Molecular DiagnosticsEdited by: Carl A.burtis, Edward R. Ashwood, David e. Bruns

Interperatation :-

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Elevated levels results from increased bilirubin production (eg hemolysis and ineffective erythropoiesis); decreased bilirubin excretion (eg; obstruction and hepatitis); and abnormal bilirubin metabolism (eg; hereditary and neonatal jaundice).conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstonesgetting into the bile ducts tumors & Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of hemolytic or pernicious anemia, transfusion reaction & a common metabolic condition termed Gilbert syndrome.

AST levels increase in viral hepatitis, blockage of the bile duct ,cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis.Ast levels may also increase after a heart attck or strenuous activity. ALT is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health. Elevated ALP levels are seen in Biliary Obstruction, Osteoblastic Bone Tumors, Osteomalacia, Hepatitis, Hyperparathyriodism, Leukemia,Lymphoma, paget's disease, Rickets, Sarcoidosis etc.

Elevated serum GGT activity can be found in diseases of the liver, Biliary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-including drugs etc.

Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum..Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom's disease. Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic - Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.

#### Renal Function Test (RFT)

Urea - SERUM

18.48

Method - Urease

15 - 39

mg/dl

		NA BHAVESH TAN	IK		Age/Sex	: 33 Year(s)	
UHID	: SHHM.5255	6			Order Date	: 12/11/202	2 10:29
Episode	: OP						
Ref. Doctor	:				Mobile No	: 969976717	
					DOB	: 09/06/198	
					Facility	SEVENHILI	LS HOSPITAL, MUMBAI
BUN - SERUM		8.64				mg/dl	4 - 18
Method - Urease	-GLDH						
Creatinine - SEF	RUM	0.63				mg/dl	0.5 - 1.1
Method - Jaffes I	Kinetic						
References:							
1)Pack Insert of	-						
2) TIETZ Textbo	ook of Clinical chem	istry and Molecular D.	iagnosticsEdited by: Ca	rl A.burtis, Edw	vard R. Ashwood, Da	vid e. Bruns	
Interpretation:-							
	-		long with the creatinine monitor people with acu		-	-	v be
used to evaluate	e a person's general		ordered as part of a ren				
Sample No : 00	<i>(CMP).</i> 0248638C	Collection Date :	12/11/22 13:11	Ack Data :	12/11/2022 14:09	Report	Date : 12/11/22 14:13
Sample No . Of	02400300	conection Date .	12/11/22 13.11	Ack Date .	12/11/2022 14.09	Report	Date . 12/11/22 14.15
GLUCOSE-PLA	SMA POST						
<u>PRANDIAL</u>							
			_				
Glucose,Post Pr	andial	107.1	7			mg/dl	70 - 140
American Diabet	tes Association Refe	erence Range :					
Post-Prandial Blo	ood Glucose:						
	: Up to 140mg/dL						
Pre-Diabetic: Diabetic	140-199 mg/dL :>200 mg/dL						
Deferences	-						
<i>References: 1)Pack Insert of</i>	Bio system						
2) TIETZ Textbo	ook of Clinical chem	istry and Molecular D	iagnostics Edited by: Ca	arl A.burtis, Edv	vard R. Ashwood,Da	vid e. Bruns	
Interpretation :-							
Conditions that of	can result in an ele	-	evel include: Acromega				,
Conditions that of stroke for instan	can result in an ele nce), Chronic kidney	v disease, Cushing syr	evel include: Acromega ndrome, Excessive cons dition characterized by	umption of foo	d, Hyperthyroidism,P	Pancreatitis.	
Conditions that o stroke for instan A low level of glu nervous system	can result in an ele nce), Chronic kidney lucose may indicate symptoms (sweatir	v disease, Cushing syr hypoglycemia, a con ng, palpitations, hung	ndrome, Excessive cons dition characterized by er, trembling, and anxie	umption of foo a drop in blood ety), then begir	d, Hyperthyroidism,F glucose to a level w hs to affect the brain	ancreatitis. here first it causes (causing confusio	5
Conditions that d stroke for instan A low level of glu nervous system hallucinations, b	can result in an elet nce), Chronic kidney lucose may indicate symptoms (sweatir lurred vision, and s	v disease, Cushing syr hypoglycemia, a con ng, palpitations, hung cometimes even coma	ndrome, Excessive cons dition characterized by a	umption of foo a drop in blood ety), then begir od glucose level	d, Hyperthyroidism,F   glucose to a level w ns to affect the brain   (hypoglycemia) maj	Pancreatitis. here first it causes (causing confusio / be	5

Patient Name	: Mrs. MEGHNA BHAVESH TANK
UHID	: SHHM.52556
Episode	: OP
Ref. Doctor	:

Age/Sex	: 33 Year(s) / Female
Order Date	: 12/11/2022 10:29
Mobile No	: 9699767177
DOB	: 09/06/1989
Facility	: SEVENHILLS HOSPITAL, MUMBAI

End of Report



Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept.

.

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

HAEMATOLOGY

Test Name	Result		Unit	Ref. Range
Sample No: 00248609A	Collection Date : 12/11/22 10:33	Ack Date : 12/11/2022 10:50	Report	Date : 12/11/22 11:06
COMPLETE BLOOD COUNT	(CBC) - EDTA WHOLE BLOOD			
Total WBC Count	9.04		x10^3/ul	4.00 - 10.00
Neutrophils	55		%	40.00 - 80.00
Lymphocytes	27.4		%	20.00 - 40.00
Eosinophils	12.4 🔺		%	1.00 - 6.00
Monocytes	5.0		%	2.00 - 10.00
Basophils	0.2 ▼		%	1.00 - 2.00
Absolute Neutrophils Count	4.98		x10^3/ul	2.00 - 7.00
Absolute Lymphocytes Count	2.48		x10^3/ul	0.80 - 4.00
Absolute Eosinophils Count	1.12 🔺		x10^3/ul	0.02 - 0.50
Absolute Monocytes Count	0.45		x10^3/ul	0.12 - 1.20
Absolute Basophils Count	0.01		x10^3/ul	0.00 - 0.10
RBCs	4.85 ⊾		x10^6/ul	3.80 - 4.80
Haemoglobin	13.8		gm/dl	12.00 - 15.00
Hematocrit	39.7 ▼		%	40.00 - 50.00

Patient Name: Mrs. MEGHNA BHAVEUHID: SHHM.52556Episode: OPRef. Doctor:	ESH TANK	Age/Sex Order Date Mobile No DOB Facility	: 96997671 : 09/06/19	22 10:29 77
MCV	82.0 ▼		fl	83.00 - 101.00
МСН	28.6		pg	27.00 - 32.00
МСНС	34.8 ▲		gm/dl	31.50 - 34.50
RED CELL DISTRIBUTION WIDTH-CV (RDW-CV)	12.9		%	11.00 - 16.00
RED CELL DISTRIBUTION WIDTH-SD (RDW-SD)	40.3		fl	35.00 - 56.00
Platelet	276		x10^3/ul	150.00 - 410.00
MPV	8.1		fl	6.78 - 13.46
PLATELET DISTRIBUTION WIDTH (PDW)	15.5		%	9.00 - 17.00
PLATELETCRIT (PCT)	0.222		%	0.11 - 0.28

NOTE: References are from "Interpretations of Diagnostic Tests" by Wallach & "Fundamentals of Clinical Chemistry" By Tietz

#### NOTE :-

The International Council for Standardization in Haematology (ICSH) recommends reporting of absolute counts of various WBC subsets for clinical decision making. This test has been performed on a fully automated 5 part differential cell counter which counts over 10,000 WBCs to derive differential counts. A complete blood count is a blood panel that gives information about the cells in a patient's blood, such as the cell count for each cell type and the concentrations of Hemoglobin and platelets. The cells that circulate in the bloodstream are generally divided into three types: white blood cells (leukocytes), red blood cells (erythrocytes), and platelets (thrombocytes). Abnormally high or low counts may be physiological or may indicate disease conditions, and hence need to be interpreted clinically.

#### ERYTHROCYTE SEDIMENTATION RATE (ESR)

22 🔺

mm/hr 0 - 20

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Method: Westergren Method

INTERPRETATION :-

ESR is a non-specific phenomenon, its measurement is clinically useful in disorders associated with an increased production of acute-phase proteins. it provides an index of progress of the disease in rheumatoid arthritis or tuberculosis, and it is of considerable value in diagnosis of temporal arteritis and polymyalgia rheumatica. It is often used if multiple myeloma is suspected, but when the myeloma is non-secretory or light chain, a normal ESR does not exclude this diagnosis.

An elevated ESR occurs as an early feature in myocardial infarction. Although a normal ESR cannot be taken to exclude the presence of organic disease, the vast majority of acute or chronic infections and most neoplastic and degenerative diseases are associated with changes in the plasma proteins that increased ES values. An increased ESR in subjects who are HIV seropositive seems to be an early predictive marker of progression toward acquired immune deficiency syndrome (AIDS).

The ESR is influenced by age, stage of the menstrual cycle and medications taken (corticosteroids, contraceptive pills). It is especially low (0–1 mm) in polycythaemia, hypofibrinogenaemia and congestive cardiac failure and when there are abnormalities of the red cells such as poikilocytosis, spherocytosis, or sickle cells. In cases of performance enhancing drug intake by athletes the ESR values are generally lower than the usual value for the individual and as a result of the increase in haemoglobin (i.e. the effect of secondary polycythaemia).

End of Report

Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept.

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

### **HISTOPATHALOGY AND CYTOLOGY**

Test Name		Resi	ult					
Sample No :	O0248626B	Collection Date :	12/11/22 11:49	Ack Date :	12/11/2022 11:56	Report Date :	12/11/22 14:23	

### ROUTINE CERVICOVAGINAL PAP SMEAR

REPORT

C-GY-162/22

#### CLINICAL DETAILS :

LMP: 28/10/2022 PS: Cervix/vagina appears healthy Minimal white discharge present

#### **MATERIAL RECEIVED :**

2 wet- fixed conventional cervico-vaginal smears received.

#### **MICROSCOPIC EXAMINATION :**

The smears are satisfactory for evaluation. Endocervical / transformation zone component is present. Benign superficial & intermediate & parabasal squamous cells noted. Few polymorphonuclear leucocytes seen. Altered bacterial flora (coccobacilli) is observed. Dysplastic cells are not seen.

### **IMPRESSION**:

Negative for intraepithelial lesion or malignancy.

NOTE :-The 2014 Bethesda system for reporting cervical cytology was followed.

#### Comments :

Cervicovaginal cytology is a screening test primarily for squamous cancer and precursors and has associated false-negative and false-positive results. Regular sampling and follow-up of unexplainded clinical signs and symptoms are recommended to minimize ffalse negative results.

End of Report



Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Dr.Nipa Dhorda MD Pathologist

.

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
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#### IMMUNOLOGY

Test Name	Result		Unit	Ref. Range
Sample No: 00248609C	Collection Date : 12/11/22 10:33	Ack Date : 12/11/2022 11:01	Report I	Date : 12/11/22 12:17
T3 - SERUM Method - CLIA	113.2		ng/dl	70.00 - 204.00
T4 - SERUM Method - CLIA	6.77		ug/dL	4.60 - 10.50
TSH - SERUM Method - CLIA	2.77		uIU/ml	0.40 - 4.50

Reference Ranges (T3) Pregnancy: First Trimester 81 - 190 Second Trimester & Third Trimester 100 - 260

Reference Ranges (TSH) Pregnancy: 1st Trimester : 0.1 – 2.5 2nd Trimester : 0.2 – 3.0 3rd Trimester : 0.3 – 3.0

#### Reference:

1.Clinical Chemistry and Molecular Diagnostics, Tietz Fundamentals, 7th Edition & Endocronology Guideliens

Interpretation :-

It is recommended that the following potential sources of variation should be considered while interpreting thyroid hormone results: 1. Thyroid hormones undergo rhythmic variation within the body this is called circadian variation in TSH secretion: Peak levels are seen between 2-4 am. Minimum levels seen between 6-10 am. This variation may be as much as 50% thus, influence of sampling time needs to be considered for clinical interpretation.

2. Circulating forms of T3 and T4 are mostly reversibly bound with Thyroxine binding globulins (TBG), and to a lesser extent with albumin and Thyroid binding PreAlbumin. Thus the conditions in which TBG and protein levels alter such as chronic liver disorders, pregnancy, excess of estrogens, androgens, anabolic steroids and glucocorticoids may cause misleading total T3, total T4 and TSH interpretations.

3. Total T3 and T4 levels are seen to have physiological rise during pregnancy and in patients on steroid treatment.

4. T4 may be normal the presence of hyperthyroidism under the following conditions : T3 thyrotoxicosis, Hypoproteinemia related reduced binding, during intake of certain drugs (eg Phenytoin, Salicylates etc)

5. Neonates and infants have higher levels of T4 due to increased concentration of TBG

6. TSH levels may be normal in central hypothyroidism, recent rapid correction of hypothyroidism or hyperthyroidism, pregnancy, phenytoin therapy etc.

7. TSH values of <0.03 uIU/mL must be clinically correlated to evaluate the presence of a rare TSH variant in certain individuals which is undetectable by conventional methods.

8. Presence of Autoimmune disorders may lead to spurious results of thyroid hormones

9. Various drugs can lead to interference in test results.

10. It is recommended that evaluation of unbound fractions, that is free T3 (fT3) and free T4 (fT4) for clinic-pathologic correlation, as these are the metabolically active forms.

Patient Name	: Mrs. MEGHNA BHAVESH TANK
UHID	: SHHM.52556
Episode	: OP
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Age/Sex	: 33 Year(s) / Female
Order Date	: 12/11/2022 10:29
Mobile No	: 9699767177
DOB	: 09/06/1989
Facility	: SEVENHILLS HOSPITAL, MUMBAI

End of Report



Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept.

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Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
Episode	: OP		
Ref. Doctor	:	Mobile No	: 9699767177
		DOB	: 09/06/1989
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Urinalysis						
Test Name	Result		Unit Ref. Range			
Sample No : 00248609D	Collection Date : 12/11/22 10:33	Ack Date : 12/11/2022 10:52	Report Date : 12/11/22 14:06			
Physical Examination						
QUANTITY	40		ml			
Colour	Pale Yellow					
Appearance	Slightly Hazy					
DEPOSIT	Absent		Absent			
рН	Acidic					
Specific Gravity	1.020					
Chemical Examination						
Protein	Absent		Absent			
Sugar	Absent		Absent			
ketones	Absent		Absent			
Occult Blood	NEGATIVE		Absent			
Bile Salt	Absent		Absent			
Bile Pigments	Absent		Absent			

Patient Name : Mrs. MEGHN UHID : SHHM.52556 Episode : OP Ref. Doctor :		Age/Sex Order Date Mobile No DOB Facility	: 33 Year(s) / Female : 12/11/2022 10:29 : 9699767177 : 09/06/1989 : SEVENHILLS HOSPITAL	, MUMBAI
Urobilinogen	NORMAL		Absent	
NITRATE	Absent			
LEUKOCYTES	Absent			
Microscopic Examination				
Puscells	6-8		/HPF	
Epithelial Cells	4-6		/HPF	
RBC	Absent		/HPF Absent	
Cast	Absent		/LPF Absent	
Crystal	Absent		/HPF Absent	
Amorphous Materials	Absent		Absent	
Yeast	Absent		Absent	
Bacteria	Absent		Absent	
<u>URINE SUGAR AND KETONE</u> (FASTING)	<u>E</u>			
Sugar	Absent			
ketones	Absent			

	Patient Name	: Mrs. MEGHNA BHAVESH TANK	Age/Sex	: 33 Year(s) / Female
	UHID	: SHHM.52556	Order Date	: 12/11/2022 10:29
	Episode	: OP		
	Ref. Doctor	:	Mobile No	: 9699767177
			DOB	: 09/06/1989
			Facility	: SEVENHILLS HOSPITAL, MUMBAI
- 1				

### URINE SUGAR AND KETONE (PP)

Sugar

Absent

ketones

.

Absent

End of Report



Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept.

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## **DIAGNOSTICS REPORT**

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Order Date	: 12/11/2022 10:29
Age/Sex	: 33 Year(s)/Female	Report Date	: 12/11/2022 13:13
UHID	: SHHM.52556	IP No	:
Ref. Doctor	: Self	Facility	: SEVENHILLS HOSPITAL, MUMBAI

#### **USG ABDOMEN**

Liver is normal in size (15 cm) and shows bright echotexture. No focal liver parenchymal lesion is seen. Intrahepatic portal and biliary radicles are normal.

Gall-bladder is physiologically distended. No evidence of intraluminal calculus is seen. Wall thickness appears normal. No evidence of peri-cholecystic fluid is seen.

Portal vein and CBD are normal in course and calibre.

Visualised part of pancreas appears normal in size and echotexture. No evidence of duct dilatation or parenchymal calcification seen.

Spleen is normal in size (11 cm) and echotexture. No focal lesion is seen in the spleen.

Right kidney measures 10.9 x 4.0 cm. Left kidney measures 10.6 x 4.7 cm.

Both the kidneys are normal in size, shape and echotexture. Cortico-medullary differentiation is maintained. No evidence of calculus or hydronephrosis on either side.

There is no free fluid in abdomen and pelvis. **IMPRESSION:** 

Grade I fatty liver.



Dr.Sagar Shriramlingam Garge, MBBS,DMRE

## **DIAGNOSTICS REPORT**

Patient Name	: Mrs. MEGHNA BHAVESH TANK	Order Date	: 12/11/2022 10:29
Age/Sex	: 33 Year(s)/Female	Report Date	: 12/11/2022 13:25
UHID Ref. Doctor	: SHHM.52556 : Self	IP No Facility	: : : SEVENHILLS HOSPITAL, MUMBAI

# X-RAY CHEST PA VIEW

Both lungs are clear.

The frontal cardiac dimensions are normal.

The pleural spaces are clear.

Both hilar shadows are normal in position and density.

No diaphragmatic abnormality is seen.

The soft tissues and bony thorax are normal.

IMPRESSION: No pleuroparenchymal lesion is seen.

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Dr.Sagar Shriramlingam Garge, MBBS,DMRE