DIAGNOSTICS REPORT

Patient Name	: Mr. PRASHANT SANWARIYA	Order Date	: 25/02/2023 09:35
Age/Sex	: 32 Year(s)/Male	Report Date	: 25/02/2023 12:18
UHID	: SHHM.59375	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) RegNo: 3393/09/2003

Patient Name	: Mr. PRASHANT SANWARIYA
UHID	: SHHM.59375
Episode	: OP
Ref. Doctor	: Self

Age/Sex	: 32 Year(s) / Male
Order Date	: 25/02/2023 09:35
Mobile No	: 9571627525
DOB	: 19/03/1990
Facility	: SEVENHILLS HOSPITAL, MUMBAI

Blood Bank

Test Name			Result					
Sample No :	O0260622A	Collection Date :	25/02/23 09:35	Ack Date :	25/02/2023 10:50	Report Date :	25/02/23 12:10	

BLOOD GROUPING/ CROSS-MATCHING BY SEMI AUTOMATION

BLOOD GROUP (ABO)	' AB '
Rh Type	POSITIVE

Method - Column Agglutination

REMARK: THE REPORTED RESULTS PERTAIN TO THE SAMPLE RECEIVED AT THE BLOOD CENTRE.

Interpretation:

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.

RegNo: 2006/03/1680

Patient Name	: Mr. PRASHANT SANWARIYA	Age/Sex	: 32 Year(s) / Male
UHID	: SHHM.59375	Order Date	: 25/02/2023 09:35
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9571627525
		DOB	: 19/03/1990
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

HAEMATOLOGY

Test Name			Result			Unit	Ref.	Range
Sample No :	O0260622A	Collection Date :	25/02/23 09:35	Ack Date :	25/02/2023 10:02		Report Date :	25/02/23 11:07
COMPLETE BLOOD COUNT (CBC) - EDTA WHOLE BLOOD								
Total WBC	Count		8	3.31			x10^3/ul	4.00 - 10.00
Neutrophils	5		Į.	50.9			%	40.00 - 80.00
Lymphocyt	es		3	38.7			%	20.00 - 40.00
Eosinophils	5		I.	5.4			%	1.00 - 6.00
Monocytes			2	1.4			%	2.00 - 10.00
Basophils			().6 ▼			%	1.00 - 2.00
Absolute N	eutrophils		2	1.23			x10^3/ul	2.00 - 7.00
Count								
Absolute Ly	ymphocytes			3.22			x10^3/ul	0.80 - 4.00
Count								
Absolute E	osinophils		().45			x10^3/ul	0.02 - 0.50
Count							40.00/ 1	0.40.4.00
	onocytes Count).36			x10^3/ul	0.12 - 1.20
	asophils Count).05			x10^3/ul	0.00 - 0.10
RBCs				5.02			x10^6/ul	4.50 - 5.50
Hemoglobi				13.7			gm/dl	13.00 - 17.00
Hematocrit	:			12.6			%	40.00 - 50.00
MCV				34.8			fl	83.00 - 101.00
MCH				27.2			pg	27.00 - 32.00
MCHC				32.1			gm/dl	31.50 - 34.50
-	DISTRIBUTION		1	12.2			%	11.00 - 16.00
WIDTH-CV			-	~ -			a	
-			2	37.7			fl	35.00 - 56.00
WIDTH-SD	(RDW-SD)			277			x10^3/ul	150.00 410.00
Platelet								150.00 - 410.00
MPV				10.9			fl 0/	6.78 - 13.46
	DISTRIBUTION		1	15.8			%	9.00 - 17.00
WIDTH (PE	-).303 ⊾			%	0.11 - 0.28
PLATELETC	CRIT (PCT)		(J.303 A			%	0.11 - 0.28

Patient Name	: Mr. PRASHANT SANWARIYA	Age/Sex	: 32 Year(s) / Male
UHID	: SHHM.59375	Order Date	: 25/02/2023 09:35
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Ref. Doctor	: Self	Mobile No	: 9571627525
		DOB	: 19/03/1990
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

NOTE: Wallach's Interpretation of Diagnostic Tests. 11th Ed, Editors: Rao LV. 2021

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) ESR

18

mm/hr 0 - 20

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 may occur 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 ESR values.

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. RegNo: 2006/03/1680

Patient Name	: Mr. PRASHANT SANWARIYA	Age/Sex	: 32 Year(s) / Male
UHID	: SHHM.59375	Order Date	: 25/02/2023 09:35
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9571627525
		DOB	: 19/03/1990
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Biochemistry								
Test Name	Result	Un	it Ref.	Range				
Sample No: 00260622A	Collection Date : 25/02/23 09:35	Ack Date : 25/02/2023 10:02	Report Date :	25/02/23 11:33				
<u>GLYCOSLYATED</u> HAEMOGLOBIN (HBA1C) HbA1c		5.78 A	%	4 to 6%				
				Non-diabetic 6.07.0% Excellent control 7.08.0% Fair to good control 8.010% Unsatisfactory control ABOVE 10% Poor control				
 HbA1c may be falsely low in diabets evaluates diabetes over 15 days. Inappropriately low HbA1c values n hypertriglyceridemia, chronic liver dise with estimation of HbA1c, causing fals HbA1c may be increased in patients Inappropriately higher values of Hb hyperbilirubinemia and large doses of Trends in HbA1c are a better indica Any sample with >15% HbA1c shot below 4% should prompt additional st HbA1c target in pregnancy is to atta HbA1c target in paediatric age groun Method : turbidimetric inhibition immute 	tic control. It reflects the mean plasma glu ics with hemolytic disease. In these individ may be reported due to hemolysis, recent l ease.Drugs like dapsone, ribavirin, antiretro rely low values. Is with polycythemia or post-splenectomy. A1c may be caused due to iron deficiency, aspirin. Itor of diabetic control than a solitary test. uld be suspected of having a hemoglobin w udies to determine the possible presence ain level <6 % .	uals a plasma fructosamine level may be u blood transfusion, acute blood loss, oviral drugs, trimethoprim, may also cause vitamin B12 deficiency, alcohol intake, ure variant, especially in a non-diabetic patient. of variant hemoglobin.	interference mia,	90 - 126				
Sample No: 00260622B	Collection Date : 25/02/23 09:35	Ack Date : 25/02/2023 10:16	Report Date :	25/02/23 11:34				
<u>GLUCOSE-PLASMA-FAST</u> ING Clusses Easting		15.72 🛦	ma/dl	70 - 110				
Glucose,Fasting	-	113.72 A	mg/dl	70 - 110				

Patient Name	: Mr. PRASHANT SANWARIYA	Age/Sex	: 32 Year(s) / Male
UHID	: SHHM.59375	Order Date	: 25/02/2023 09:35
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9571627525
		DOB	: 19/03/1990
		Facility	: SEVENHILLS HOSPITAL, MUMBAI
1			

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, 6th Ed, Editors: Rifai et al. 2018

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 :	O0260622C	Collection Date :	25/02/23 09:35	Ack Date :	25/02/2023 10:15	Report Date :	25/02/23 13:08

Lipid Profile Total Cholesterol	238.26	mg/dl	Reference Values :
			Up to 200 mg/dL -
			Desirable
			200-239 mg/dL - Borderline HIgh
			>240 mg/dL - High
Triglycerides	191.5	mg/dl	Reference Values:
rigiyeendes			Up to 150 mg/dL -
			Normal
			150-199 mg/dL -
			Borderline High
			200-499 mg/dL -
			High
			>500 mg/dL - Very High
Method - Enzymatic			ngn
HDL Cholesterol	44.43	mg/dl	0 - 60
Method - Enzymatic immuno inhibition		•	
LDL Cholesterol	155.53 🔺	mg/dl	0 - 130
Method - Calculated			
VLDL Cholesterol	38.30	mg/dl	0 - 40
Method - Calculated			
Total Cholesterol / HDL	5.36 ▲	RATIO	0 - 5
Cholesterol Ratio -			
Calculated			

UHID : Episode : Ref. Doctor :	Mr. PRASHANT SANWARIYA SHHM.59375 OP Self		er Date : 25/0 ile No : 9571 : 19/1	ear(s) / Male 2/2023 09:35 .627525 03/1990 ENHILLS HOSP	ITAL, MUMBAI
Method - Calculated LDL / HDL Choles Ratio - Calculated Method - Calculated References: 1)Pack Insert of Bio s 2) Tietz Textbook Of	d	3.50 iditors: Rifai et al. 2018		RATIO	0 - 4.3
adults. Triglycerides c hours after eating. Ev different days are not 2. HDL-Cholesterol: H tissues and carries it t increased risk of hear HDL cholesterol value risk factor. 3. LDL-Cholesterol: D acceptable. Values be levels of LDL cholester inflammation, or cirrh		much as 5 to 10 times higher re, modest changes in fasting d" cholesterol, because it rem for men and less than 50 mg ding the LDL-C level. The NC ated as a negative I risk factors. For young adult Values greater than 160 mg/o	r than fasting levels ju triglycerides measure loves excess choleste /dL for women, there EP guidelines suggest ts, less than 120 mg/ dL are considered higu	ist a few ed on rol from is an that an dL is h. Low	
Uric Acid (Seru Uric Acid Method - Uricase References: 1)Pack Insert of Bio s 2) TIETZ Textbook o		8.6 ▲ Carl A.burtis,Edward R. Ashv	vood,David e. Bruns	mg/dl	3.50 - 7.20
including our DNA. In inflammation and pair	by the breakdown of purines. Purines are nitrogen-cont creased concentrations of uric acid can cause crystals to n characteristic of gout. Low values can be associated w to toxic compounds, and rarely as the result of an inher. Test (o form in the joints, which ca with some kinds of liver or kid	n lead to the joint ney diseases, Fancon	i.	
SGOT (Aspartate Transaminase) - <i>Method - IFCC</i>		43.45 ⊾		U/L	0 - 35
SGPT (Alanine Transaminase) - <i>Method - IFCC</i>	SERUM	96.29 🛦		U/L	0 - 45
Total Bilirubin - S Method - Diazo	SERUM	0.36		mg/dl	0 - 2
Direct Bilirubin - Method - Diazotizatio		0.15		mg/dl	0 - 0.4

Patient Name UHID Episode Ref. Doctor	: Mr. PRASHANT SANWARIYA : SHHM.59375 : OP : Self		Age/Sex Order Date Mobile No DOB Facility	: 32 Year(s) / Male : 25/02/2023 09:35 : 9571627525 : 19/03/1990 : SEVENHILLS HOSF	
Indirect Biliru Calculated		0.21		mg/dl	0.1 - 0.8
Method - Calcula Alkaline Phos SERUM	sphatase -	52.99		U/L	0 - 115
Method - IFCC A Total Protein Method - Biuret	- SERUM	6.96		gm/dl	6 - 7.8
Albumin - SE Method - Bromo	RUM o Cresol Green(BCG)	4.47		gm/dl	3.5 - 5.2
Globulin - Ca Method - Calcula		2.49		gm/dl	2 - 4
A:G Ratio Method - Calcula	ated	1.80		:1	1 - 3
Gamma Gluta Transferase (Gglutamyl ca nitroanilide - <i>Method - G gluta</i>	(GGT) - rboxy	61.01 ▲		U/L	0 - 55

References:

1)Pack Insert of Bio system

2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018

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 (

<u>RFT)</u>

Urea - SERUM

mg/dl 15 - 39

~							
Patient Name	: Mr. PRASH	ANT SANWARIYA			Age/Sex	: 32 Year(s) / Male	:
UHID	: SHHM.5937	'5			Order Date	: 25/02/2023 09:3	5
Episode	: OP					-,-,-	
Ref. Doctor	: Self				Mobile No	: 9571627525	
					DOB	: 19/03/1990	
					Facility	: SEVENHILLS HOS	SPITAL MUMBAT
					ruenty		
Mothed Ukonso							
Method - Urease BUN - SERUM			c	9.26		mg/dl	4 - 18
Method - Urease-			-			ing/ai	1 10
Creatinine - SI			().8		mg/dl	0.5 - 1.3
Method - Jaffes K	-					5.	
References:							
1)Pack Insert of B	-	try And Molecular Diag	apostics 6th Ed Edita	rc: Difai at al 2	019		
2) HELZ TEXLOOK	or climical chemis	iy And Molecular Diag		<i>13. Milai et al.</i> 20	010		
Interpretation:-							
	-	is primarily used, alor ney disease, and to mo	-		-	-	
	a person's general i						
Sample No : O	0260654B	Collection Date :	25/02/23 12:28	Ack Date :	25/02/2023 13:01	Report Date :	25/02/23 14:04
<u>GLUCOSE-PL</u> <u>PRANDIAL</u>	ASMA POST						
Glucose,Post F	Drandial		5	209.36 🛦		mg/dl	70.00 - 140.00
-	s Association Refer	ence Range :	_				
Post-Prandial Blood Glucose: Non- Diabetic: Up to 140mg/dL Pre-Diabetic: 140-199 mg/dL Diabetic :>200 mg/dL							
References:							
1)Pack Insert of B 2) Tietz Textbook	-	try And Molecular Diag	anostics, 6th Ed. Edito	rs: Rifai et al. 2	018		
2) //012 /0/100000		.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	<i></i>				
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.							
Severe neart fallui	ге, сптопискіапеу	(renar) ranure, INSUIIN	overause, rumors tha				
0 (End of Rep	DOIT		

Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept. RegNo: 2006/03/1680

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Page 5 of 5

Patient Name: Mr. PRASHANT SANWARIYAUHID: SHHM.59375Episode: OPRef. Doctor: Self

Age/Sex: 32 Year(s) / MaleOrder Date: 25/02/2023 09:35Mobile No: 9571627525DOB: 19/03/1990Facility: SEVENHILLS HOSPITAL, MUMBAI

IMMUNOLOGY

Test Name		Result			Unit	Ref.	Range
Sample No: 00260622C	Collection Date :	25/02/23 09:35	Ack Date :	25/02/2023 10:15		Report Date :	25/02/23 11:28
T3 - SERUM		1	11.4			ng/dl	70.00 - 204.00
<i>Method - CLIA</i> T4 - SERUM		c	.45			ug/dL	4.60 - 10.50
Method - CLIA		-	. 15			ug/uL	1.00 10.50
TSH - SERUM		1	.83			uIU/ml	0.40 - 4.50
Method - CLIA							
Reference Ranges (T3) Pregnancy:							
First Trimester 81 - 190							
Second Trimester & Third Trimester	r 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.



End of Report

Patient Name	: Mr. PRASHANT SANWARIYA	Age/Sex	: 32 Year(s) / Male
UHID	: SHHM.59375	Order Date	: 25/02/2023 09:35
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9571627525
		DOB	: 19/03/1990
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept. RegNo: 2006/03/1680

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Patient Name	: Mr. PRASHANT SANWARIYA	Age/Sex	: 32 Year(s) / Male
UHID	: SHHM.59375	Order Date	: 25/02/2023 09:35
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9571627525
		DOB	: 19/03/1990
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

				Urinalysis					
Test Name			Result			Unit	Ref.	Range	
Sample No :	O0260625D	Collection Date :	25/02/23 09:46	Ack Date :	25/02/2023 10:45	F	Report Date :	25/02/23 15:08	
URINE S	UGAR AND								
<u>KETONE</u>	(FASTING)								
Sugar				Absent					
ketones				Absent					
Sample No :	O0260657D	Collection Date :	25/02/23 12:34	Ack Date :	25/02/2023 13:57	F	Report Date :	25/02/23 15:08	
	UGAR AND								
KETONE									
Sugar				Absent					
ketones				Absent					
				End of Rep	ort				
	Dipa								
MD	l ipa Dhorda ologist								

DIAGNOSTICS REPORT

Patient Name	: Mr. PRASHANT SANWARIYA	Order Date	: 25/02/2023 09:35
Age/Sex	: 32 Year(s)/Male	Report Date	: 25/02/2023 12:42
UHID	: SHHM.59375	IP No	:
Ref. Doctor	: Self	Facility	SEVENHILLS HOSPITAL, MUMBAI

USG ABDOMEN

Liver is normal in size (13.8 cm) and shows bright echotexture. No focal liver parenchymal lesion is seen. Intrahepatic portal and biliary radicles are normal.

Gall-bladder is partially collapsed

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 (8.0 cm) and echotexture. No focal lesion is seen in the spleen.

Right kidney measures 8.9 x 4.7 cm. Left kidney measures 9.3 x 5.2 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.

Urinary bladder is partially distended and appears normal. No evidence of intra-luminal calculus or mass lesion.

Prostate appears grossly normal in size and echotexture.

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

Grade I fatty liver.

Alani Dr-Shubham Asrani

Dr.Shubham Asrani , MBBS, MD

RegNo: 2020/01/0042

DIAGNOSTICS REPORT

Patient Name	: Mr. PRASHANT SANWARIYA	Order Date	: 25/02/2023 09:35
Age/Sex	: 32 Year(s)/Male	Report Date	: 25/02/2023 12:22
UHID	: SHHM.59375	IP No	:
Ref. Doctor	: Self	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.

MPRESSION: No pleuroparenchymal lesion is seen.

Kula

Dr.Bhujang Pai, MBBS, MD

Consultant