DIAGNOSTICS REPORT

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Order Date	: 11/02/2023 09:27
Age/Sex	: 35 Year(s)/Male	Report Date	: 11/02/2023 10:44
UHID	: SHHM.58330	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.

Grade I 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. KIRAN SITARAM KALAMBE	
UHID	: SHHM.58330	(
Episode	: OP	
Ref. Doctor	: Self	I
		I

Age/Sex	: 35 Year(s) / Male
Order Date	: 11/02/2023 09:27
Mobile No	: 9833669094
DOB	: 18/06/1987
Facility	: SEVENHILLS HOSPITAL, MUMBAI

Blood Bank Test Name Result O0258708A 11/02/23 09:38 Ack Date : 11/02/2023 12:03 11/02/23 12:24 Sample No : Collection Date : Report Date : BLOOD GROUPING/ CROSS-MATCHING BY SEMI AUTOMATION BLOOD GROUP (ABO) 'A' NEGATIVE Rh Type Du TEST NEGATIVE Comment **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. KIRAN SITARAM KALAMBE	Age/Sex	: 35 Year(s) / Male
UHID	: SHHM.58330	Order Date	: 11/02/2023 09:27
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9833669094
		DOB	: 18/06/1987
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Biochemistry						
Test Name	Result	Unit Ref	. Range			
Sample No: 00258708A	Collection Date : 11/02/23 09:38 Ack Date :	11/02/2023 10:42 Report Date :	11/02/23 12:04			
<u>GLYCOSLYATED</u> HAEMOGLOBIN (HBA1C)						
HbA1c Method - BIOCHEMISTRY	5.21	%	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			
Estimated Average Glucose (eAG) Method - Calculated NOTES :- 1. HbA1c is used for monitoring diabe 2. HbA1c may be falsely low in diabet evaluates diabetes over 15 days. 3. Inappropriately low HbA1c values in hypertriglyceridemia, chronic liver disc with estimation of HbA1c, causing fals 4. HbA1c may be increased in patients 5. Inappropriately higher values of Hb hyperbilirubinemia and large doses of 6. Trends in HbA1c are a better indica 7. Any sample with >15% HbA1c sho below 4% should prompt additional si 8. HbA1c target in pregnancy is to att 9. HbA1c target in paediatric age grou Method : turbidimetric inhibition immu	with polycythemia or post-splenectomy. A1c may be caused due to iron deficiency, vitamin B12 de aspirin. tor of diabetic control than a solitary test. Ild be suspected of having a hemoglobin variant, especiall udies to determine the possible presence of variant hemog ain level <6 % .	uctosamine level may be used which n, acute blood loss, nethoprim, may also cause interference ficiency, alcohol intake, uremia, y in a non-diabetic patient. Similarly,	90 - 126			
Sample No: 00258708B	Collection Date : 11/02/23 09:38 Ack Date :	11/02/2023 10:47 Report Date :	11/02/23 11:12			
GLUCOSE-PLASMA-FAST ING Glucose,Fasting	94.3	mg/dl	70 - 110			

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Age/Sex	: 35 Year(s) / Male
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		DOB	: 18/06/1987
		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:

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 : 00258708C Collection Date : 11/02/23 09:38 Ack Date : 11/02/2023 10:51 Report Date : 11/02/23 12:04	
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Lipid Profile			
Total Cholesterol	143.09	mg/dl	Reference Values :
			Up to 200 mg/dL -
			Desirable
			200-239 mg/dL -
			Borderline HIgh
	83.7		>240 mg/dL - High Reference Values:
Triglycerides	83.7	mg/dl	Up to 150 mg/dL -
			Normal
			150-199 mg/dL -
			Borderline High
			200-499 mg/dL -
			High
			>500 mg/dL - Very
			High
Method - Enzymatic	64.18 🛦		0 (0
HDL Cholesterol	04.18 🛦	mg/dl	0 - 60
Method - Enzymatic immuno inhibition	62.17	mg/dl	0 - 130
LDL Cholesterol Method - Calculated	02.17	mg/ui	0 - 130
VLDL Cholesterol	16.74	mg/dl	0 - 40
Method - Calculated	10071	iiig/ ai	0 10
Total Cholesterol / HDL	2.23	RATIO	0 - 5
Cholesterol Ratio -			
Calculated			
Calculated			

¹⁾Pack Insert of Bio system

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Age/Sex	• : 35 Year(s) / Ma	e
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-		Order D	ate : 11/02/2023 09:	27
Episode	: OP			
Ref. Doctor	: Self	Mobile I		
		DOB	: 18/06/1987	
		Facility	: SEVENHILLS HC	SPITAL, MUMBAI
Method - Calculat	ted			
LDL / HDL Ch	olesterol	0.97	RATIO	0 - 4.3
Ratio - Calcula	ated			
Method - Calculat	ted			
References:				
1)Pack Insert of E 2) Tietz Texthool	810 system k Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, E	ditors: Rifai et al. 2018		
2) 11612 16210001				
Interpretation				
	When triglycerides are very high greater than 1000 mg/dL, ti			
	les change dramatically in response to meals, increasing as g. Even fasting levels vary considerably day to day. Therefor	-		
-	not considered to be abnormal.			
2. HDL-Cholester	ol: HDL- C is considered to be beneficial, the so-called "good	d" cholesterol, because it removes	excess cholesterol from	
	s it to the liver for disposal. If HDL-C is less than 40 mg/dL	-		
	heart disease that is independent of other risk factors, inclu alue greater than 60 mg/dL is protective and should be trea		idelines suggest that an	
risk factor.	and greater than of mg/ul is protective and should be trea	aleu as a negative		
	ol: Desired goals for LDL-C levels change based on individua	n risk factors. For young adults, le	ss than 120 mg/dL is	
acceptable. Value	es between 120-159 mg/dL are considered Borderline high.	Values greater than 160 mg/dL are	e considered high. Low	
	lesterol may be seen in people with an inherited lipoprotein	deficiency and in people with hyp	erthyroidism, infection,	
inflammation, or o				
Uric Acid	<u>21011)</u>	4.3	mg/dl	3.5 - 7.2
Method - Uricase		1.5	iiig/ai	5.5 7.2
References:				
1)Pack Insert of E	3io system			
2) TIETZ Textboo	ok of Clinical chemistry and Molecular DiagnosticsEdited by:	Carl A.burtis,Edward R. Ashwood	David e. Bruns	
Interpretation:-				
	ced by the breakdown of purines. Purines are nitrogen-cont	aining compounds found in the ce	ells of the body,	
	A. Increased concentrations of uric acid can cause crystals to			
	pain characteristic of gout. Low values can be associated w	,		
, , ,	ure to toxic compounds, and rarely as the result of an inher	ited metabolic defect (Wilson disea	ase).	
<u>Liver Function LFT)</u>	<u>on Test (</u>			
SGOT (Aspart	ate	20.69	U/L	0 - 35
Transaminase		_0.07	0/2	0 00
Method - IFCC) SERGIN			
SGPT (Alanine		25.04	U/L	0 - 45
Transaminase				
Method - IFCC				
Total Bilirubin	- SERLIM	0.96	mg/dl	0 - 2
Method - Diazo	JEROM		119/01	
Direct Bilirubi	n SERLIM	0.46 🔺	mg/dl	0 - 0.4
Method - Diazotiz			ing/u	0 0.1
metriou - Dia2002				

Patient Name	e : Mr. KIRAN SITARAM KALAMBE		Age/Sex	: 35 Year(s) / Male	
UHID	: SHHM.58330		Order Date	: 11/02/2023 09:27	
Episode	: OP				
Ref. Doctor	: Self		Mobile No	: 9833669094	
			DOB	: 18/06/1987	
			Facility	: SEVENHILLS HOSI	PITAL, MUMBAI
Indirect Biliru	ubin -	0.50		mg/dl	0.1 - 0.8
Calculated				-	
Method - Calcula	ated				
Alkaline Phos	sphatase -	108.66		U/L	0 - 115
SERUM					
Method - IFCC A		8.14 ▲		am /dl	6 7 9
Total Protein Method - Biuret		0.14 🛦		gm/dl	6 - 7.8
Albumin - SE		5.15		gm/dl	3.5 - 5.2
	Cresol Green(BCG)	5115		gnijai	5.5 5.2
Globulin - Ca		2.99		gm/dl	2 - 4
Method - Calcula				C .	
A:G Ratio		1.72		:1	1 - 3
Method - Calcula	ated				
Gamma Gluta	amyl	27.04		U/L	0 - 55
Transferase					
Gglutamyl ca	-				
nitroanilide -					
-	amyl carboxy nitroanilide				
Poforoncoci					

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

Dationt Name			DE		• 1-		
Patient Name	: Mr. KIRAN	SITARAM KALAM	BE		Age/Sex	: 35 Year(s) / Male	
UHID	: SHHM.5833	30			Order Date	: 11/02/2023 09:27	,
Episode	: OP						
Ref. Doctor	: Self				Mobile No	:9833669094	
					DOB	: 18/06/1987	
					Facility	: SEVENHILLS HOS	PITAL, MUMBAI
Method - Urease BUN - SERUM				7.93		mg/dl	4 - 18
Method - Urease-							
Creatinine - S				0.75		mg/dl	0.5 - 1.3
Method - Jaffes k	-					-	
References:							
1)Pack Insert of E 2) Tietz Textbook		try And Molecular Dia	gnostics, 6th Ed, Edite	ors: Rifai et al. 20	018		
Tadaman di di							
Interpretation:- The blood urea ni	itroaen or RI IN tect	t is primarily used, alo	na with the creatining	e test, to evaluate	kidnev function in a	wide range of	
	-	ney disease, and to m	-		-	-	
used to evaluate a	a person's general i	health status.					
Sample No: O	0258754B	Collection Date :	11/02/23 12:30	Ack Date :	11/02/2023 13:37	Report Date :	11/02/23 14:35
Post-Prandial Bloc	es Association Refer	rence Range :		88.7		mg/dl	70.00 - 140.00
Pre-Diabetic: 1 Diabetic	40-199 mg/dL :>200 mg/dL						
References: 1)Pack Insert of E 2) Tietz Textbook		try And Molecular Dia	gnostics, 6th Ed, Edite	ors: Rifai et al. 20	018		
stroke for instance A low level of glue nervous system sj	e), Chronic kidney (cose may indicate h ymptoms (sweating	ated blood glucose lev disease, Cushing sync hypoglycemia, a condi g, palpitations, hungen metimes even coma a	frome, Excessive cons ition characterized by r, trembling, and anxi	sumption of food, a drop in blood g ety), then begins	Hyperthyroidism,Par glucose to a level whe to affect the brain (c	ocreatitis. rre first it causes ausing confusion,	
seen with:Adrena		king excessive alcoho (renal) failure, Insulir					
seen with:Adrena		-			in (insulinomas),Starv		

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

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Patient Name	: Mr. KIRAN SITARAM KALAMBE	Age/Sex	: 35 Year(s) / Male
UHID	: SHHM.58330	Order Date	: 11/02/2023 09:27
Episode	: OP		
Ref. Doctor	: Self	Mobile No	: 9833669094
		DOB	: 18/06/1987
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

HAEMATOLOGY

Test Name			Result			Unit	Ref.	Range	
Sample No : O	0258708A	Collection Date :	11/02/23 09:38	Ack Date :	11/02/2023 10:42		Report Date :	11/02/23 11:04	
COMPLETE B	COMPLETE BLOOD COUNT (CBC) - EDTA WHOLE BLOOD								
Total WBC Cou	unt			5.66			x10^3/ul	4.00 - 10.00	
Neutrophils				69.4			%	40.00 - 80.00	
Lymphocytes				21.8			%	20.00 - 40.00	
Eosinophils				1.1			%	1.00 - 6.00	
Monocytes				7.4			%	2.00 - 10.00	
Basophils				0.3 🔻			%	1.00 - 2.00	
Absolute Neut	rophils			3.93			x10^3/ul	2.00 - 7.00	
Count									
Absolute Lymp	ohocytes			1.24			x10^3/ul	0.80 - 4.00	
Count									
Absolute Eosin	nophils			0.06			x10^3/ul	0.02 - 0.50	
Count									
Absolute Mono	•			0.42			x10^3/ul	0.12 - 1.20	
Absolute Baso	phils Count			0.01			x10^3/ul	0.00 - 0.10	
RBCs				5.80 🔺			x10^6/ul	4.50 - 5.50	
Hemoglobin				16.6			gm/dl	13.00 - 17.00	
Hematocrit				50.1 ▲			%	40.00 - 50.00	
MCV				86.5			fl	83.00 - 101.00	
MCH				28.7			pg	27.00 - 32.00	
MCHC				33.2			gm/dl	31.50 - 34.50	
RED CELL DIS WIDTH-CV (RI				12.4			%	11.00 - 16.00	
RED CELL DIS WIDTH-SD (RI	TRIBUTION			39.2			fl	35.00 - 56.00	
Platelet	עכ-זייט)			234			x10^3/ul	150.00 - 410.00	
MPV				8.4			fl	6.78 - 13.46	
				15.6			%	9.00 - 17.00	
PLATELET DIS WIDTH (PDW)				13.0			70	5.00 - 17.00	
PLATELETCRI	T (PCT)			0.197			%	0.11 - 0.28	

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Age/Sex	: 35 Year(s) / Male
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		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

5

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 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.

RegNo: 2006/03/1680

Patient Name	: Mr. KIRAN SITARAM KALAMBE
UHID	: SHHM.58330
Episode	: OP
Ref. Doctor	: Self

Age/Sex	: 35 Year(s) / Male
Order Date	: 11/02/2023 09:27
Mobile No	: 9833669094
DOB	: 18/06/1987
Facility	: SEVENHILLS HOSPITAL, MUMBAI

IMMUNOLOGY

Test Name		Result			Unit	Ref.	Range
Sample No: 00258708C	Collection Date :	11/02/23 09:38	Ack Date :	11/02/2023 10:51		Report Date :	11/02/23 12:05
T3 - SERUM Method - CLIA		:	112.5			ng/dl	70 - 204
T4 - SERUM Method - CLIA		9	9.55			ug/dL	4.6 - 10.5
TSH - SERUM		2	2.43			uIU/ml	0.4 - 4.5
Method - CLIA 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.



End of Report

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Age/Sex	: 35 Year(s) / Male
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Dr.Ritesh Kharche MD, PGD HOD, Laboratory Medicine Dept.

RegNo: 2006/03/1680

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Age/Sex	: 35 Year(s) / Male
UHID	: SHHM.58330	Order Date	: 11/02/2023 09:27
Episode	: OP		
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		DOB	: 18/06/1987
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Urinalysis								
Test Name			Result			Unit	Ref.	Range
Sample No :	O0258708D	Collection Date :	11/02/23 09:38	Ack Date :	11/02/2023 11:01		Report Date :	11/02/23 12:05
URINE SL	JGAR AND							
<u>KETONE (</u>	FASTING)							
Sugar				Absent				
ketones				Absent				
Sample No :	O0258759D	Collection Date :	11/02/23 12:38	Ack Date :	11/02/2023 12:53		Report Date :	11/02/23 14:36
 <u>URINE SL</u>	JGAR AND							
<u>KETONE (</u>	<u>(PP)</u>							
Sugar				Absent				
ketones				Absent				
				End of Rep	ort			
8	flah							

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

RegNo: 2006/03/1680

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DIAGNOSTICS REPORT

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Order Date	: 11/02/2023 09:27
Age/Sex	: 35 Year(s)/Male	Report Date	: 11/02/2023 15:17
UHID	: SHHM.58330	IP No	:
Ref. Doctor	: Self	Facility	: SEVENHILLS HOSPITAL, MUMBAI

USG ABDOMEN & PELVIS

Liver is normal in size (13.2 cm) and 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 (9.0 cm) and echotexture. No focal lesion is seen in the spleen.

Right kidney measures 9.0 x 4.1 cm. Left kidney measures 9.6 x 5.1 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 well distended and appears normal. No evidence of intra-luminal calculus or mass lesion.

Prostate appears normal in size and echotexture. It measures 3.9 x 3.0 x 2.8 cm corresponding to 17.8 cc.

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

'No significant abnormality is detected.

Dr.Rashmi Randive , MBBS, MD

DIAGNOSTICS REPORT

Patient Name	: Mr. KIRAN SITARAM KALAMBE	Order Date	: 11/02/2023 09:27
Age/Sex	: 35 Year(s)/Male	Report Date	: 11/02/2023 15:09
UHID	: SHHM.58330	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.

IMPRESSION: No pleuroparenchymal lesion is seen.

Dr.Rashmi Randive , MBBS, MD