







Patient Name : SANTUJIT SARKAR

**Age** : 38 Y 0 M 0 D

Gender : M

Lab Add. : Newtown,Kolkata-700156

Ref Dr. : Dr.MEDICAL OFFICER

Collection Date : 05/Jul/2024 10:19AM

Report Date : 05/Jul/2024 04:37PM

#### DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
SGOT/AST , GEL SERUM (Method:Modified IFCC)	21	13-40	U/L
CHLORIDE,BLOOD (Method:ISE INDIRECT)	107	99-109	mEq/L
PHOSPHORUS-INORGANIC,BLOOD (Method:Phosphomolybdate/UV)	2.7	2.4-5.1 mg/dL	mg/dL
ALKALINE PHOSPHATASE (Method:IFCC standardization )	93	46-116	U/L
BILIRUBIN (DIRECT) (Method:Vanadate oxidation)	0.20	<0.2	mg/dL
BILIRUBIN (TOTAL), GEL SERUM			
BILIRUBIN (TOTAL) (Method:Vanadate oxidation)	0.70	0.3-1.2	mg/dL
POTASSIUM,BLOOD (Method:ISE INDIRECT)	4.00	3.5-5.5	mEq/L
UREA,BLOOD (Method:Urease with GLDH)	19.3	19-49	mg/dL
CREATININE, BLOOD (Method:Jaffe, alkaline picrate, kinetic)	0.79	0.7-1.3	mg/dL
GLUCOSE,FASTING (Method:Gluc Oxidase Trinder)	95	Impaired Fasting-100-125 .~Diabetes- >= 126.~Fasting is defined as no caloric intake for at least 8 hours.	mg/dL

In the absence of unequivocal hyperglycemia, diagnosis requires two abnormal test results from the same sample or in two separate test samples.

#### Reference :

ADA Standards of Medical Care in Diabetes – 2020. Diabetes Care Volume 43, Supplement 1.

CALCIUM,BLOOD (Method:Arsenazo III)	9.80	8.7-10.4	mg/dL
URIC ACID,BLOOD (Method:Uricase/Peroxidase)	5.20	3.5-7.2	mg/dL
SGPT/ALT (Method:Modified IFCC)	24	7-40	U/L
SODIUM,BLOOD (Method:ISE INDIRECT)	142	132 - 146	mEq/L

\*\*\* End Of Report \*\*\*









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Report Date : 05/Jul/2024 04:37PM



Test Name Result Bio Ref. Interval Unit

Dr Neepa Chowdhury MBBS, MD(Biochemistry) SECTION DIRECTOR AND SENIOR CONSULTANT BIOCHEMIST Reg no. WBMC 62456









 Patient Name
 : SANTUJIT SARKAR
 Ref Dr.
 : Dr.MEDICAL OFFICER

 Age
 : 38 Y 0 M 0 D
 Collection Date
 : 05/Jul/2024 10:19AM

 Gender
 : M
 Report Date
 : 05/Jul/2024 08:14PM



#### DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit	
				_
THYROID PANEL (T3, T4, TSH), GEL SERUM				
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	1.02	0.60-1.81 ng/ml	ng/ml	
T4-TOTAL (THYROXINE) (Method:CLIA)	9.8	3.2-12.6	μg/dL	
TSH (THYROID STIMULATING HORMONE) (Method:CLIA)	1.916	0.55-4.78	μlU/mL	

Serum TSH levels exhibit a diurnal variation with the peak occurring during the night and the nadir, which approximates to 50% of the peak value, occurring between 1000 and 1600 hours.[1,2]

#### References:

- 1. Bugalho MJ, Domingues RS, Pinto AC, Garrao A, Catarino AL, Ferreira T, Limbert E and Sobrinho L. Detection of thyroglobulin mRNA transcripts in peripheral blood of
- individuals with and without thyroid glands: evidence for thyroglobulin expression by blood cells. Eur J Endocrinol 2001;145:409-13.
- 2. Bellantone R, Lombardi CP, Bossola M, Ferrante A, Princi P, Boscherini M et al. Validity of thyroglobulin mRNA assay in peripheral blood of postoperative thyroid carcinoma patients in predicting tumor recurrence varies according to the histologic type: results of a prospective study. Cancer 2001;92:2273-9.

#### **BIOLOGICAL REFERENCE INTERVAL**: [ONLY FOR PREGNANT MOTHERS]

Trimester specific TSH LEVELS during pregnancy: FIRST TRIMESTER:  $0.10-3.00~\mu$  IU/mL SECOND TRIMESTER: 0.20 -3.50  $\mu$  IU/mL THIRD TRIMESTER: 0.30 -3.50  $\mu$  IU/mL

#### **References:**

- 1. Erik K. Alexander, Elizabeth N. Pearce, Gregory A. Brent, Rosalind S. Brown, Herbert Chen, Chrysoula Dosiou, William A. Grobman, Peter Laurberg, John H. Lazarus, Susan J. Mandel, Robin P. Peeters, and Scott Sullivan. Thyroid.Mar 2017.315-389. <a href="https://doi.org/10.1089/thy.2016.0457">https://doi.org/10.1089/thy.2016.0457</a>
  2. Kalza S. Agarwal S. Agarwal S. Agarwal R. Randhir S. Trimester, specific thyroid-stimulating hormone: An indian perspective. Indian J. Endocr Metab.
- 2. Kalra S, Agarwal S, Aggarwal R, Ranabir S. Trimester-specific thyroid-stimulating hormone: An indian perspective. Indian J Endocr Metab 2018;22:1-4.

\*\*\* End Of Report \*\*\*

Dr. SANCHAYAN SINHA MBBS, MD, DNB (BIOCHEMISTRY) CONSULTANT BIOCHEMIST Reg No. WBMC 63214

**Lab No.**: DUN/05-07-2024/SR9329508 Page 3 of 14









Lab No. : DUN/05-07-2024/SR9329508 Lab Add. : Newtown, Kolkata-700156

**Patient Name** : SANTUJIT SARKAR Ref Dr. : Dr.MEDICAL OFFICER :38 Y 0 M 0 D **Collection Date** : 05/Jul/2024 10:19AM Age Gender

Report Date : 05/Jul/2024 05:19PM



#### DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit

GLYCATED HAEMOGLOBIN (HBA1C), EDTA WHOLE BLOOD

GLYCATED HEMOGLOBIN (HBA1C) 4.9 \*\*\*FOR BIOLOGICAL REFERENCE %

> INTERVAL DETAILS, PLEASE REFER TO THE BELOW MENTIONED REMARKS/NOTE WITH ADDITIONAL CLINICAL

**INFORMATION \*\*\*** 

HbA1c (IFCC) 30.0 mmol/mol

(Method:HPLC)

Clinical Information and Laboratory clinical interpretation on Biological Reference Interval:

Low risk / Normal / non-diabetic : <5.7% (NGSP) / < 39 mmol/mol (IFCC) Pre-diabetes/High risk of Diabetes: 5.7%-6.4% (NGSP) / 39 - < 48 mmol/mol (IFCC) : >/= 6.5% (NGSP) / > 48 mmol/mol (IFCC) Diabetics-HbA1c level

Analyzer used :- Bio-Rad-VARIANT TURBO 2.0

Method: HPLC Cation Exchange

#### Recommendations for glycemic targets

- Ø Patients should use self-monitoring of blood glucose (SMBG) and HbA1c levels to assess glycemic control.
- Ø The timing and frequency of SMBG should be tailored based on patients' individual treatment, needs, and goals.
- Ø Patients should undergo HbA1c testing at least twice a year if they are meeting treatment goals and have stable glycemic control.
- Ø If a patient changes treatment plans or does not meet his or her glycemic goals, HbA1c testing should be done quarterly.
- Ø For most adults who are not pregnant, HbA1c levels should be <7% to help reduce microvascular complications and macrovascular disease. Action suggested >8% as it indicates poor control.
- Ø Some patients may benefit from HbA1c goals that are stringent.

Result alterations in the estimation has been established in many circumstances, such as after acute/ chronic blood loss, for example, after surgery, blood transfusions, hemolytic anemia, or high erythrocyte turnover; vitamin B<sub>12</sub>/ folate deficiency, presence of chronic renal or liver disease; after administration of high-dose vitamin E / C; or erythropoietin treatment.

Reference: Glycated hemoglobin monitoring BMJ 2006; 333;586-8

#### References

- Chamberlain JJ, Rhinehart AS, Shaefer CF, et al. Diagnosis and management of diabetes: synopsis of the 2016 American Diabetes Association Standards of Medical Care in Diabetes. Ann Intern Med. Published online
- 1 March 2016. doi:10.7326/M15-3016.

  Mosca A, Goodall I, Hoshino T, Jeppsson JO, John WG, Little RR, Miedema K, Myers GL, Reinauer H, Sacks DB, Weykamp CW. International Federation of Clinical Chemistry and Laboratory Medicine, IFCC Scientific Division. Global standardization of glycated hemoglobin measurement: the position of the IFCC Working Group. Clin Chem Lab Med. 2007;45(8):1077-1080.

#### PDF Attached

TOTAL PROTEIN [BLOOD] ALB:	LO RATIO , .			
TOTAL PROTEIN (Method:BIURET METHOD)	<u>8.50</u>	5.7-8.2 g/dL	g/dL	
ALBUMIN (Method:BCG Dye Binding)	<u>5.0</u>	3.2-4.8 g/dL	g/dL	
GLOBULIN (Method:Calculated)	<u>3.50</u>	1.8-3.2	g/dl	
AG Ratio (Method:Calculated)	1.43	1.0-2.5		

URIC ACID, URINE, SPOT URINE			
URIC ACID, SPOT URINE (Method:URICASE)	57.00	37-92 mg/dL	mg/dL

(Method:URICASE)				
LIPID PROFILE, GEL SERUM				
CHOLESTEROL-TOTAL (Method:Enzymatic)	223	Desirable: < 200 mg/dL Borderline high: 200-239 mg/dL High: > or =240 mg/dL	mg/dL	
TRIGLYCERIDES	133	Normal:: < 150,	mg/dL	

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**Lab No.** : DUN/05-07-2024/SR9329508 L

: SANTUJIT SARKAR

**Age** : 38 Y 0 M 0 D

**Patient Name** 

Gender : M

Lab Add. : Newtown,Kolkata-700156

Ref Dr. : Dr.MEDICAL OFFICER

Report Date : 05/Jul/2024 05:19PM

: 05/Jul/2024 10:19AM

**Collection Date** 



#### DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
(Method:GPO-Trinder)		BorderlineHigh::150-199, High:: 200-499, VeryHigh::>500	
HDL CHOLESTEROL (Method:Elimination/catalase)	44	< 40 - Low 40-59- Optimum 60 - High	mg/dl
LDL CHOLESTEROL DIRECT (Method:Elimination / Catalase)	147	OPTIMAL: <100 mg/dL, Near optimal/ above optimal: 100- 129 mg/dL, Borderline high: 130-159 mg/dL, High: 160-189 mg/dL, Very high: >=190 mg/dL	mg/dL
VLDL (Method:Calculated)	32	< 40 mg/dl	mg/dl
CHOL HDL Ratio (Method:Calculated)	5.1	LOW RISK 3.3-4.4 AVERAGE RISK 4.47-7.1 MODERATE RISK 7.1-11.0 HIGH RISK >11.0	

Reference: National Cholesterol Education Program. Executive summary of the third report of The National Cholesterol Education Program (NCEP) Expert Panel on detection, evaluation, and treatment of high blood cholesterol in adults (Adult Treatment Panel III). JAMA. May 16 2001;285(19):2486-97.

\*\*\* End Of Report \*\*\*

DR. ANANNYA GHOSH MBBS, MD (Biochemistry) Consultant Biochemist Reg No. WBMC 73007

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Unit

Lab No. : DUN/05-07-2024/SR9329508 Lab Add. : Newtown, Kolkata-700156

Result

**Patient Name** : SANTUJIT SARKAR Ref Dr. : Dr.MEDICAL OFFICER :38 Y 0 M 0 D **Collection Date** : 05/Jul/2024 01:37PM Age Gender

: 05/Jul/2024 05:50PM Report Date



#### DEPARTMENT OF BIOCHEMISTRY

92*		mg/dL
	92*	92* Impaired Glucose Tolerance-140 to 199. Diabetes>= 200.

Bio Ref. Interval

\*NOTE: The lower value of Plasma Glucose (PP) compared to that of Plasma Glucose(F), may be interpreted having due to regard to the history of the case with particular reference to Diabetes, if any including the time and dose of antidiabetic drug administered, if any.

Blood glucose level is maintained by a very complex integrated mechanism involving critical interplay of release of hormones and action of enzymes on key metabolic pathways resulting in a smooth transition normally from a high level of glucose influx following meal / glucose intake to a basal  $level\ after\ 2-3\ hrs.\ or\ so.\ Excluding\ a limentary\ hypoglycemia,\ renal\ glycosuria,\ hereditary\ fructose\ intolerance\ and\ Galactosemia,\ the\ possible$ causes of post prandial reactive hypoglycemia (PRH) include high insulin sensitivity, exaggerated response of insulin and glucagon like peptide 1, defects in counter-regulation, very lean and /or anxious individuals, after massive weight reduction etc.

The test should be performed as described by the WHO, using a glucose load containing the equivalent of 75-g anhydrous glucose dissolved in water.

In the absence of unequivocal hyperglycemia, diagnosis requires two abnormal test results from the same sample or in two separate test samples.

**Test Name** 

ADA Standards of Medical Care in Diabetes - 2020. Diabetes Care Volume 43, Supplement 1.

\*\*\* End Of Report \*\*\*

Dr. Sudeshna Bara M.B.B.S MD. (Biochemistry) (Consultant Biochemist) Reg No. WBMC 64124

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 Patient Name
 : SANTUJIT SARKAR
 Ref Dr.
 : Dr.MEDICAL OFFICER

 Age
 : 38 Y 0 M 0 D
 Collection Date
 : 05/Jul/2024 10:19AM

 Gender
 : M
 Report Date
 : 05/Jul/2024 05:32PM



#### DEPARTMENT OF HAEMATOLOGY

Test Name Result Bio Ref. Interval Unit

#### BLOOD GROUP ABO+RH [GEL METHOD], EDTA WHOLE BLOOD

ABO A

(Method:Gel Card)

RH POSITIVE

(Method:Gel Card)

#### **TECHNOLOGY USED: GEL METHOD**

#### ADVANTAGES:

- $\cdot$   $\;$  GeI card allows simultaneous forward and reverse grouping.
- · Card is scanned and record is preserved for future reference.
- · Allows identification of Bombay blood group.
- Daily quality controls are run allowing accurate monitoring.

#### Historical records check not performed.

CBC WITH PLATELET (THROMBOCYTE)	COUNT, EDTA WHOLE BLO	OD	
HEMOGLOBIN (Method:PHOTOMETRIC)	13.6	13 - 17	g/dL
WBC	7.8	4 - 10	*10^3/µL
(Method:DC detection method) RBC	4.55	4.5 - 5.5	*10^6/µL
(Method:DC detection method)	4.00	4.5 - 5.5	10 0/μΕ
PLATELET (THROMBOCYTE) COUNT (Method:DC detection method/Microscopy)	174	150 - 450*10^3	*10^3/µL
DIFFERENTIAL COUNT			
NEUTROPHILS	60	40 - 80 %	%
(Method:Flowcytometry/Microscopy) LYMPHOCYTES	30	20 - 40 %	%
(Method:Flowcytometry/Microscopy)	30	20 - 40 %	/0
MONOCYTES	08	2 - 10 %	%
(Method:Flowcytometry/Microscopy) EOSINOPHILS	02	1 - 6 %	%
(Method:Flowcytometry/Microscopy)	02	1 - 0 %	70
BASOPHILS	00	0-0.9%	%
(Method:Flowcytometry/Microscopy)			
CBC SUBGROUP	00.4	40. 50.0/	0/
HEMATOCRIT / PCV (Method:Calculated)	<u>39.1</u>	40 - 50 %	%
MCV	86.0	83 - 101 fl	fl
(Method:Calculated)			
MCH (Method:Calculated)	29.9	27 - 32 pg	pg
MCHC	<u>34.8</u>	31.5-34.5 gm/dl	gm/dl
(Method:Calculated)		•	
RDW - RED CELL DISTRIBUTION WIDTH (Method:Calculated)	<u>14.7</u>	11.6-14%	%
PDW-PLATELET DISTRIBUTION WIDTH	33.3	8.3 - 25 fL	fL
(Method:Calculated)		<del>-</del>	
MPV-MEAN PLATELET VOLUME	13.0	7.5 - 11.5 fl	
(Method:Calculated)			

ESR (ERYTHROCYTE SEDIMENTATION RATE), EDTA WHOLE BLOOD

1stHour 06 0.00 - 20.00 mm/hr mm/hr

(Method:Westergren)

**Lab No.** : DUN/05-07-2024/SR9329508

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Patient Name : SANTUJIT SARKAR

**Age** : 38 Y 0 M 0 D

Gender : M

Lab Add. : Newtown,Kolkata-700156

Ref Dr. : Dr.MEDICAL OFFICER

Collection Date : 05/Jul/2024 10:19AM

Report Date : 05/Jul/2024 05:32PM

#### DEPARTMENT OF HAEMATOLOGY

Test Name Result Bio Ref. Interval Unit

\*\*\* End Of Report \*\*\*

Dr. KAUSHIK DEY
MD (PATHOLOGY)
CONSULTANT PATHOLOGIST

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Reg No. WBMC 66405



Patient Name : SANTUJIT SARKAR Ref Dr. : Dr.MEDICAL OFFICER

Age : 38 Y 0 M 0 D Collection Date

**Gender** : M Report Date : 05/Jul/2024 12:51PM



#### DEPARTMENT OF X-RAY

### X-RAY CHEST PA VIEW

Lab Add.

Bilateral lung fields appear normal.

Bilateral costophrenic angles are unremarkable.

Bilateral hila and vascular markings are unremarkable.

Domes of diaphragm are normal in morphology and contour.

Cardiac size is within normal limits.

Bony thoracic cage appears normal.

#### IMPRESSION:

No obvious abnormality detected.

No evidence of fracture or dislocation.

Recommended clinical correlation.

\*\*\* End Of Report \*\*\*

Dr. Manish Kumar Jha MD Radiodiagnosis

**Lab No.** : DUN/05-07-2024/SR9329508 Page 9 of 14









 Patient Name
 : SANTUJIT SARKAR
 Ref Dr.
 : Dr.MEDICAL OFFICER

 Age
 : 38 Y 0 M 0 D
 Collection Date
 : 06/Jul/2024 07:23AM

 Gender
 : M
 Report Date
 : 06/Jul/2024 11:40AM



#### DEPARTMENT OF CLINICAL PATHOLOGY

Test Name Result Bio Ref. Interval Unit

JRINE ROUTINE ALL, ALL , URINE PHYSICAL EXAMINATION				
COLOUR	PALE YELLOW			
APPEARANCE	SLIGHTLY HAZY			
CHEMICAL EXAMINATION				
pH (Method:Dipstick (triple indicator method))	6.0	4.6 - 8.0		
SPECIFIC GRAVITY (Method:Dipstick (ion concentration method))	1.020	1.005 - 1.030		
PROTEIN (Method:Dipstick (protein error of pH dicators)/Manual)	NOT DETECTED	NOT DETECTED		
GLUCOSE (Method:Dipstick(glucose-oxidase-peroxidase	NOT DETECTED	NOT DETECTED		
nethod)/Manual) KETONES (ACETOACETIC ACID, ACETONE) (Method:Dipstick (Legals test)/Manual)	NOT DETECTED	NOT DETECTED		
BLOOD (Method:Dipstick (pseudoperoxidase reaction))	NOT DETECTED	NOT DETECTED		
BILIRUBIN (Method:Dipstick (azo-diazo reaction)/Manual)	NEGATIVE	NEGATIVE		
UROBILINOGEN (Method:Dipstick (diazonium ion reaction)/Manual)	NEGATIVE	NEGATIVE		
NITRITE (Method:Dipstick (Griess test))	NEGATIVE	NEGATIVE		
LEUCOCYTE ESTERASE (Method:Dipstick (ester hydrolysis reaction))  MICROSCOPIC EXAMINATION	NEGATIVE	NEGATIVE		
LEUKOCYTES (PUS CELLS) (Method:Microscopy)	0-1	0-5	/hpf	
EPITHELIAL CELLS (Method:Microscopy)	1-2	0-5	/hpf	
RED BLOOD CELLS (Method:Microscopy)	NOT DETECTED	0-2	/hpf	
CAST (Method:Microscopy)	NOT DETECTED	NOT DETECTED		
CRYSTALS (Method:Microscopy)	NOT DETECTED	NOT DETECTED		
BACTERIA (Method:Microscopy)	SCANTY	NOT DETECTED		
YEAST (Method:Microscopy)	NOT DETECTED	NOT DETECTED		

#### Note:

- $1. \ All \ urine \ samples \ are \ checked \ for \ adequacy \ and \ suitability \ before \ examination.$
- 2. Analysis by urine analyzer of dipstick is based on reflectance photometry principle. Abnormal results of chemical examinations are confirmed by manual methods.
- 3. The first voided morning clean-catch midstream urine sample is the specimen of choice for chemical and microscopic analysis.
- 4. Negative nitrite test does not exclude urinary tract infections.
- 5. Trace proteinuria can be seen in many physiological conditions like exercise, pregnancy, prolonged recumbency etc.
- 6. False positive results for glucose, protein, nitrite, urobilinogen, bilirubin can occur due to use of certain drugs, therapeutic dyes, ascorbic acid, cleaning agents used in urine collection container.
- 7. Discrepancy between results of leukocyte esterase and blood obtained by chemical methods with corresponding pus cell and red blood cell count by microscopy can occur due to cell lysis.
- 8. Contamination from perineum and vaginal discharge should be avoided during collection, which may falsely elevate epithelial cell count and show presence of bacteria

  Lab No. : DUN/05-07-2024/SR9329508 Page 10 of 14









**Patient Name** : SANTUJIT SARKAR

Age :38 Y 0 M 0 D

Gender : M Lab Add. : Newtown, Kolkata-700156

Ref Dr. : Dr.MEDICAL OFFICER

**Collection Date** : 06/Jul/2024 07:23AM

Report Date : 06/Jul/2024 11:40AM

#### DEPARTMENT OF CLINICAL PATHOLOGY

Bio Ref. Interval **Test Name** Result Unit

and/or yeast in the urine.

\*\*\* End Of Report \*\*\*

Kaushik Dr. KAUSHIK DEY MD (PATHOLOGY) CONSULTANT PATHOLOGIST

Reg No. WBMC 66405



Patient Name : SANTUJIT SARKAR Ref Dr. : Dr.MEDICAL OFFICER

Age : 38 Y 0 M 0 D Collection Date

**Gender** : M Report Date : 05/Jul/2024 04:04PM



#### DEPARTMENT OF CARDIOLOGY

Lab Add.

		DEFARTMENT OF CARDIC	
		E.C.G. REPORT	
DATA HEART RATE	61	Bpm	
PR INTERVAL	142	Ms	
QRS DURATION	94	Ms	
QT INTERVAL	390	Ms	
QTC INTERVAL	393	Ms	
AXIS P WAVE	52	Degree	
QRS WAVE	59	Degree	
T WAVE	26	Degree	
IMPRESSION		Normal sinus rhythm, within normal limits.	

\*\*\* End Of Report \*\*\*

Dr. KAUSIK PAL MD DM (Card) Reg No-WBMC-56578



Patient Name : SANTUJIT SARKAR Ref Dr. : Dr.MEDICAL OFFICER

Age : 38 Y 0 M 0 D Collection Date :

**Gender** : M Report Date : 05/Jul/2024 03:57PM

#### DEPARTMENT OF ULTRASONOGRAPHY

#### **DEPARTMENT OF ULTRASONOGRAPHY**

#### REPORT ON EXAMINATION OF WHOLE ABDOMEN

**LIVER:** It is normal in size (13 cm), normal in shape and parenchymal echopattern. No focal lesion of altered echogenicity is seen. Intrahepatic biliary radicles are not dilated. The portal vein branches and hepatic veins are normal.

**GALL BLADDER:** Well distended lumen shows no inralumnial calculus or mass. Wall thickness is normal. No pericholecystic collection is noted.

**PORTA HEPATIS:** The portal vein (0.70 cm) is normal in caliber with clear lumen. The common bile duct is normal in caliber. Visualized lumen is clear till visualised extent. Common bile duct measures approx 0.25 cm in diameter. *Extreme lower end of common bile duct is not visualised due to bowel gas shadow.* 

<u>PANCREAS</u>: It is normal in shape, size and echopattern. Main pancreatic duct is not dilated. No focal lesion of altered echogenicity is seen. The peripancreatic region shows no abnormal fluid collection.

**SPLEEN**: It is normal in shape, size (9.7 cm) and shows homogeneous echopattern. No focal lesion is seen. No abnormal venous dilatation is seen in the splenic hilum.

**KIDNEYS**: Both Kidneys are normal in shape, size and position. Cortical echogenicity and thickness are normal with normal cortico-medullary differentiation in both kidneys. No calculus, hydronephrosis is noted in either side. The perinephric region shows no abnormal fluid collection.

RIGHT KIDNEY measures 9.8 cm LEFT KIDNEY measures 9.4 cm

**URETER**: Both ureters are not dilated. No calculus is noted in either side.

**PERITONEUM & RETROPERITONEUM**: The aorta and IVC are normal. Lymph nodes are not enlarged. No free fluid is seen in peritoneum.

**URINARY BLADDER:** It is adequately distended providing optimum scanning window. The lumen is clear and wall thickness is normal.

**PROSTATE**: It is normal in shape, size and echopattern. No focal lesion is seen. Capsule is smooth.

Prostate measures: 2.8 x 4.0 x 3.2 cm. Weight 19 gms.

IMPRESSION:

No significant abnormality detected.

Please correlate clinically.

#### Kindly note

- Ultrasound is not the modality of choice to rule out subtle bowel lesion.
- Please Intimate us for any typing mistakes and send the report for correction within 7 days.
- The science of Radiological diagnosis is based on the interpretation of various shadows produced by both the normal and abnormal tissues and are not always conclusive. Further biochemical and radiological investigation & clinical correlation is required to enable the clinician to reach the final diagnosis.

The report and films are not valid for medico-legal purpose.

**Lab No.**: DUN/05-07-2024/SR9329508 Page 13 of 14



Patient Name : SANTUJIT SARKAR Ref Dr. : Dr.MEDICAL OFFICER

Age : 38 Y 0 M 0 D Collection Date

Gender : M Report Date : 05/Jul/2024 03:57PM



## DEPARTMENT OF ULTRASONOGRAPHY <u>Patient Identity not verified</u>

Lab Add.

DR. NAMRATA CHATTERJEE MBBS,CONSULTANT SONOLOGIST

Reg No: 79092

## SURAKSHA DIAGNOSTIC,RAJARHAT,KOLKATA BIO-RAD VARIANT-II TURBO CDM5.4. SN-16122

# PATIENT REPORT V2TURBO A1c 2.0

Patient Data Analysis Data

Sample ID: D02135744166 Analysis Performed: 05/JUL/2024 14:42:01

Patient ID: SR9329508 Injection Number: 3034
Name: SANTUJIT SARKAR Run Number: 42
Physician: Rack ID: 0003

Sex: M Tube Number: 7

DOB: Report Generated: 05/JUL/2024 14:50:45

Operator ID: ANUP

Comments:

	NGSP		Retention	Peak
Peak Name	%	Area %	Time (min)	Area
Unknown		0.1	0.114	3321
A1a		0.8	0.163	22688
A1b		1.2	0.226	33850
F		0.7	0.276	19922
LA1c		1.8	0.392	50812
A1c	4.9		0.491	126340
P3		3.3	0.781	95354
P4		1.2	0.858	34322
Ao		86.7	1.001	2516422

Total Area: 2,903,030

#### HbA1c (NGSP) = 4.9 % HbA1c (IFCC) = 30 mmol/mol

