

**Lab No.** : SRE/08-04-2023/SR7503529  
**Patient Name** : SUDEEP BHATTACHARYYA  
**Age** : 31 Y 0 M 0 D  
**Gender** : M

**Lab Add.** : Newtown, Kolkata-700156  
**Ref Dr.** : Dr.MEDICAL OFFICER  
**Collection Date:** 08/Apr/2023 08:19AM  
**Report Date** : 08/Apr/2023 12:07PM



Test Name	Result	Unit	Bio Ref. Interval	Method
<b>SODIUM, BLOOD , GEL SERUM</b>				
SODIUM,BLOOD	141	mEq/L	132 - 146 mEq/L	ISE INDIRECT
<b>POTASSIUM, BLOOD , GEL SERUM</b>				
POTASSIUM,BLOOD	3.90	mEq/L	3.5-5.5 mEq/L	ISE INDIRECT
<b>GLUCOSE, FASTING , BLOOD, NAF PLASMA</b>				
GLUCOSE,FASTING	87	mg/dL	Impaired Fasting-100-125 ~Diabetes- >= 126.~Fasting is defined as no caloric intake for at least 8 hours.	Gluc Oxidase Trinder

*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.

**\*CHLORIDE, BLOOD , .**

CHLORIDE,BLOOD 105 mEq/L 99-109 mEq/L ISE INDIRECT

□

Dr NEEPA CHOWDHURY  
MBBS MD (Biochemistry)  
Consultant Biochemist



Lab No. : SR7503529 Name : SUDEEP BHATTACHARYYA Age/G : 31 Y 0 M 0 D / M Date : 08-04-2023

**PHOSPHORUS-INORGANIC, BLOOD , GEL SERUM**

PHOSPHORUS-INORGANIC,BLOOD 3.9 mg/dL 2.4-5.1 mg/dL Phosphomolybdate/UV

**BILIRUBIN (DIRECT) , GEL SERUM**

BILIRUBIN (DIRECT) 0.10 mg/dL <0.2 mg/dL Vanadate oxidation

**SGOT/AST , GEL SERUM**

SGOT/AST 46 U/L 13-40 U/L Modified IFCC

**UREA,BLOOD , GEL SERUM**

UREA,BLOOD 27.8 mg/dL 19-49 mg/dL Urease with GLDH

**URIC ACID, BLOOD , GEL SERUM**

URIC ACID,BLOOD 8.00 mg/dL 3.5-7.2 mg/dL Uricase/Peroxidase

**SUGGESTED FOLLOW-UP**

**URIC ACID, URINE, SPOT URINE**

URIC ACID, SPOT URINE 68.00 mg/dL 37-92 mg/dL URICASE

**THYROID PANEL (T3, T4, TSH) , GEL SERUM**

T3-TOTAL (TRI IODOTHYRONINE) 1.26 ng/ml 0.60-1.81 ng/ml CLIA

T4-TOTAL (THYROXINE) 11.0 µg/dL 3.2-12.6 µg/dL CLIA

TSH (THYROID STIMULATING HORMONE) 3.09 µIU/mL 0.55-4.78 µIU/mL CLIA

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 µ IU/mL

SECOND TRIMESTER: 0.20 -3.50 µ IU/mL

THIRD TRIMESTER : 0.30 -3.50 µ 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. <http://doi.org/10.1089/thy.2016.0457>
2. Kalra S, Agarwal S, Aggarwal R, Ranabir S. Trimester-specific thyroid-stimulating hormone: An indian perspective.



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Indian J Endocr Metab 2018;22:1-4.

**SGPT/ALT , GEL SERUM**

SGPT/ALT **81** U/L 7-40 U/L Modified IFCC

**CALCIUM, BLOOD**

CALCIUM,BLOOD 9.30 mg/dL 8.7-10.4 mg/dL Arsenazo III

**BILIRUBIN (TOTAL) , GEL SERUM**

BILIRUBIN (TOTAL) 0.30 mg/dL 0.3-1.2 mg/dL Vanadate oxidation

**ALKALINE PHOSPHATASE , GEL SERUM**

ALKALINE PHOSPHATASE 92 U/L 46-116 U/L IFCC standardization

**CREATININE, BLOOD**

0.84 mg/dL 0.7-1.3 mg/dL Jaffe, alkaline picrate, kinetic

**TOTAL PROTEIN [BLOOD] ALB:GLO RATIO , .**

TOTAL PROTEIN 7.50 g/dL 5.7-8.2 g/dL BIURET METHOD

ALBUMIN 4.5 g/dL 3.2-4.8 g/dL BCG Dye Binding

GLOBULIN 3.00 g/dl 1.8-3.2 g/dl Calculated

AG Ratio 1.50 1.0 - 2.5 Calculated

**LIPID PROFILE , GEL SERUM**

CHOLESTEROL-TOTAL 197 mg/dL Desirable: < 200 mg/dL Enzymatic  
Borderline high: 200-239 mg/dL  
High: > or =240 mg/dL

TRIGLYCERIDES **256** mg/dL Normal:: < 150, GPO-Trinder  
BorderlineHigh::150-199,  
High:: 200-499,  
VeryHigh::>500

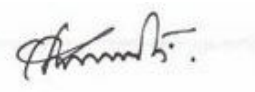
HDL CHOLESTEROL **34** mg/dl < 40 - Low Elimination/catalase  
40-59- Optimum  
60 - High

LDL CHOLESTEROL DIRECT **141** mg/dL OPTIMAL : <100 mg/dL, Elimination / Catalase  
Near optimal/ above optimal :  
100-129 mg/dL,  
Borderline high : 130-159 mg/dL,  
High : 160-189 mg/dL,  
Very high : >=190 mg/dL

VLDL 22 mg/dl < 40 mg/dl Calculated

CHOL HDL Ratio 5.8 LOW RISK 3.3-4.4 AVERAGE Calculated  
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.

  
**Dr. SUPARBA CHAKRABARTI**  
MBBS, MD(BIOCHEMISTRY)  
Consultant Biochemist



Lab No. : SR7503529      Name : SUDEEP BHATTACHARYYA      Age/G : 31 Y 0 M 0 D / M      Date : 08-04-2023

**CBC WITH PLATELET (THROMBOCYTE) COUNT , EDTA WHOLE BLOOD**

HEMOGLOBIN	13.5	g/dL	13 - 17	PHOTOMETRIC
WBC	10.0	*10 <sup>3</sup> /μL	4 - 10	DC detection method
RBC	<b>4.45</b>	*10 <sup>6</sup> /μL	4.5 - 5.5	DC detection method
PLATELET (THROMBOCYTE) COUNT	186	*10 <sup>3</sup> /μL	150 - 450*10 <sup>3</sup> /μL	DC detection method/Microscopy

**DIFFERENTIAL COUNT**

NEUTROPHILS	47	%	40 - 80 %	Flowcytometry/Microscopy
LYMPHOCYTES	40	%	20 - 40 %	Flowcytometry/Microscopy
MONOCYTES	08	%	2 - 10 %	Flowcytometry/Microscopy
EOSINOPHILS	04	%	1 - 6 %	Flowcytometry/Microscopy
BASOPHILS	<b>01</b>	%	0-0.9%	Flowcytometry/Microscopy

**CBC SUBGROUP**

HEMATOCRIT / PCV	41.0	%	40 - 50 %	Calculated
MCV	92.1	fl	83 - 101 fl	Calculated
MCH	30.5	pg	27 - 32 pg	Calculated
MCHC	33.1	gm/dl	31.5-34.5 gm/dl	Calculated
RDW - RED CELL DISTRIBUTION WIDTH	<b>15.1</b>	%	11.6-14%	Calculated
PDW-PLATELET DISTRIBUTION WIDTH	31.7	fL	8.3 - 25 fL	Calculated
MPV-MEAN PLATELET VOLUME	13.1		7.5 - 11.5 fl	Calculated

**ESR (ERYTHROCYTE SEDIMENTATION RATE) , EDTA WHOLE BLOOD**

1stHour	17	mm/hr	0.00 - 20.00 mm/hr	Westergren
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**BLOOD GROUP ABO+RH [GEL METHOD] , EDTA WHOLE BLOOD**

ABO	A	Gel Card
RH	POSITIVE	Gel Card

**TECHNOLOGY USED: GEL METHOD**

**ADVANTAGES :**

- Gel 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.**

**DR. NEHA GUPTA**  
**MD, DNB (Pathology)**  
**Consultant Pathologist**



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**URINE ROUTINE ALL, ALL , URINE**

**PHYSICAL EXAMINATION**

COLOUR PALE YELLOW  
APPEARANCE SLIGHTLY HAZY

**CHEMICAL EXAMINATION**

pH	5.0	4.6 - 8.0	Dipstick (triple indicator method)
SPECIFIC GRAVITY	1.020	1.005 - 1.030	Dipstick (ion concentration method)
PROTEIN	NOT DETECTED	NOT DETECTED	Dipstick (protein error of pH indicators)/Manual
GLUCOSE	NOT DETECTED	NOT DETECTED	Dipstick (glucose-oxidase-peroxidase method)/Manual
KETONES (ACETOACETIC ACID, ACETONE)	NOT DETECTED	NOT DETECTED	Dipstick (Legals test)/Manual
BLOOD	NOT DETECTED	NOT DETECTED	Dipstick (pseudoperoxidase reaction)
BILIRUBIN	NEGATIVE	NEGATIVE	Dipstick (azo-diazo reaction)/Manual
UROBILINOGEN	NEGATIVE	NEGATIVE	Dipstick (diazonium ion reaction)/Manual
NITRITE	NEGATIVE	NEGATIVE	Dipstick (Griess test)
LEUCOCYTE ESTERASE	NEGATIVE	NEGATIVE	Dipstick (ester hydrolysis reaction)

**MICROSCOPIC EXAMINATION**

LEUKOCYTES (PUS CELLS)	0-1	/hpf	0-5	Microscopy
EPITHELIAL CELLS	1-2	/hpf	0-5	Microscopy
RED BLOOD CELLS	NOT DETECTED	/hpf	0-2	Microscopy
CAST	NOT DETECTED		NOT DETECTED	Microscopy
CRYSTALS	NOT DETECTED		NOT DETECTED	Microscopy
BACTERIA	NOT DETECTED		NOT DETECTED	Microscopy
YEAST	NOT DETECTED		NOT DETECTED	Microscopy

**Note:**

- All urine samples are checked for adequacy and suitability before examination.
- Analysis by urine analyzer of dipstick is based on reflectance photometry principle. Abnormal results of chemical examinations are confirmed by manual methods.
- The first voided morning clean-catch midstream urine sample is the specimen of choice for chemical and microscopic analysis.
- Negative nitrite test does not exclude urinary tract infections.
- Trace proteinuria can be seen in many physiological conditions like exercise, pregnancy, prolonged recumbency etc.
- 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.
- 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.
- Contamination from perineum and vaginal discharge should be avoided during collection, which may falsely elevate epithelial cell count and show presence of bacteria and/or yeast in the urine.

**Dr. PANKTI PATEL**  
**MBBS , MD (PATHOLOGY)**  
**CONSULTANT PATHOLOGIST**



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[PDF Attached](#)

**GLYCATED HAEMOGLOBIN (HBA1C) , EDTA WHOLE BLOOD**

GLYCATED HEMOGLOBIN (HBA1C)	5.4	%	***FOR BIOLOGICAL REFERENCE INTERVAL DETAILS , PLEASE REFER TO THE BELOW MENTIONED REMARKS/NOTE WITH ADDITIONAL CLINICAL INFORMATION ***
HbA1c (IFCC)	36.0	mmol/mol	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)  
 Diabetics-HbA1c level : >/= 6.5% (NGSP) / > 48 mmol/mol (IFCC)

**Analyzer used : Bio-Rad-VARIANT TURBO 2.0**  
**Method : HPLC Cation Exchange**

**Recommendations for glycemc targets**

- Ø Patients should use self-monitoring of blood glucose (SMBG) and HbA1c levels to assess glycemc 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 glycemc control.
- Ø If a patient changes treatment plans or does not meet his or her glycemc 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:**

1. 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.
2. 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.

**GLUCOSE, PP , BLOOD, NAF PLASMA**

GLUCOSE,PP	121	mg/dL	Impaired Glucose Tolerance-140 Gluc Oxidase Trinder to 199. Diabetes>= 200.
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**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.**

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

**DR. ANANNYA GHOSH**  
**MBBS, MD (Biochemistry)**  
**Consultant Biochemist**



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**Collection Date:**  
**Report Date** : 08/Apr/2023 12:33PM




**X-RAY REPORT OF CHEST (PA)**

**FINDINGS :**

No active lung parenchymal lesion is seen.  
Both the hila are normal in size, density and position.  
Mediastinum is in central position. Trachea is in midline.  
Domes of diaphragm are smoothly outlined. Position is within normal limits.  
Lateral costo-phrenic angles are clear.  
The cardio-thoracic ratio is normal.  
Bony thorax reveals no definite abnormality.

**IMPRESSION :**

**Normal study.**

  
Dr. Anoop Sastry  
MBBS, DMRT(CAL)  
CONSULTANT RADIOLOGIST  
Registration No.: WB-36628

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Patient Name : SUDEEP BHATTACHARYYA  
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Gender : M

Lab Add. :  
Ref Dr. : Dr.MEDICAL OFFICER  
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Report Date : 08/Apr/2023 03:50PM



**DEPARTMENT OF CARDIOLOGY**  
**REPORT OF E.C.G.**

DATA		
HEART RATE	76	Bpm
PR INTERVAL	158	Ms
QRS DURATION	90	Ms
QT INTERVAL	364	Ms
QTC INTERVAL	409	Ms
AXIS		
P WAVE	- 90	Degree
QRS WAVE	27	Degree
T WAVE	24	Degree
<b>IMPRESSION</b>	<b>:</b>	<b>Sinus rhythm.</b>
		<b>Normal axis.</b>
		<b>No significant ischemic changes.</b>
		<b>Please correlate clinically.</b>

**DR. SUBHASISH BERA**  
MBBS (Cal), PGDCC  
Reg. No: 59285(WBMC)



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**Report Date** : 08/Apr/2023 12:13PM



**DEPARTMENT OF ULTRASONOGRAPHY**  
**REPORT ON EXAMINATION OF WHOLE ABDOMEN**

**LIVER**

**Liver is enlarged in size (18.79 cm), having grade II fatty changes.** No focal parenchymal lesion is evident. Intrahepatic biliary radicles are not dilated. Branches of portal vein are normal.

**PORTA**

The appearance of porta is normal. Common bile duct is normal (0.40 cm) with no intraluminal pathology (calculi /mass) could be detected at its visualised part. Portal vein is normal (1.00 cm) at porta.

**GALLBLADDER**

Gallbladder is distended. Wall thickness appears normal. No intraluminal pathology (calculi/mass) could be detected. Sonographic Murphys sign is negative.

**PANCREAS**

**Fatty infiltration of pancreas noted.** No Calcular disease noted. Pancreatic duct is not dilated. No peri-pancreatic collection of fluid noted.

**SPLEEN**

Spleen is normal in size (10.74 cm). Homogenous and smooth echotexture without any focal lesion. Splenic vein at hilum appears normal. No definite collaterals could be detected.

**KIDNEYS**

Both kidneys are normal in shape, size (Rt. kidney 11.20 cm. & Lt. kidney 11.23 cm) axes & position. Cortical echogenicity appears normal maintaining cortico-medullary differentiation. Margin is regular and cortical thickness is uniform. No calcular disease noted. No hydronephrotic changes detected.

**URETERS**

Visualised part of upper ureters are not dilated.

**URINARY BLADDER**

Urinary bladder is distended, wall thickness appeared normal. No intraluminal pathology (calculi / mass) could be detected.

**PROSTATE**

Prostate is normal in size. Echotexture appears within normal limits. No focal alteration of its echogenicity could be detectable.

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It measures : 3.09 cm. x 3.06 cm. x 2.70 cm.

Approximate weight could be around = 13.36 gms.

**RETROPERITONEUM & PERITONEUM**

No ascites noted. No definite evidence of any mass lesion detected. No detectable evidence of enlarged lymph nodes noted. Visualized part of aorta & IVC are within normal limit.

**IMPRESSION :**

- 1) Hepatomegaly with grade II fatty changes.**
- 2) Fatty infiltrations of pancreas.**

**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.*

*Patient Identity not verified.*

**DR. S. K. MONDAL**  
MBBS, CBET  
(Sonologist)

**Patient Data**

Sample ID: D02135119676  
 Patient ID: SR7503529  
 Name:  
 Physician:  
 Sex:  
 DOB:

**Analysis Data**

Analysis Performed: 08/APR/2023 13:00:27  
 Injection Number: 2007U  
 Run Number: 47  
 Rack ID: 0002  
 Tube Number: 7  
 Report Generated: 08/APR/2023 13:09:34  
 Operator ID: ASIT

Comments:

Peak Name	NGSP %	Area %	Retention Time (min)	Peak Area
A1a	---	0.9	0.156	21932
A1b	---	1.1	0.216	26179
F	---	0.7	0.265	17334
LA1c	---	1.8	0.395	43124
A1c	5.4	---	0.500	106592
P3	---	3.4	0.790	82431
P4	---	1.2	0.867	28774
Ao	---	86.4	0.984	2069349

Total Area: 2,395,714

**HbA1c (NGSP) = 5.4 %**      HbA1c (IFCC) = 36 mmol/mol

