



Lab No.	: MDG/08-06-2024/SR9211871	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: TANMOY GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 41 Y 9 M 27 D	Collection Date	: 08/Jun/2024 09:20AM
Gender	: M	Report Date	: 08/Jun/2024 12:35PM

**DEPARTMENT OF BIOCHEMISTRY**

Test Name	Result	Bio Ref. Interval	Unit
BILIRUBIN (DIRECT) , GEL SERUM (Method:Vanadate oxidation)	0.10	<0.2	mg/dL
SGOT/AST (Method:Modified IFCC)	24	13-40	U/L
SODIUM,BLOOD (Method:ISE INDIRECT)	142	132 - 146	mEq/L
CREATININE, BLOOD (Method:Jaffe, alkaline picrate, kinetic)	0.81	0.7-1.3	mg/dL
GLUCOSE,FASTING (Method:Gluc Oxidase Trinder)	91	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.

PHOSPHORUS-INORGANIC,BLOOD (Method:Phosphomolybdate/UV)	3.0	2.4-5.1 mg/dL	mg/dL
GLUCOSE,PP (Method:Gluc Oxidase Trinder)	104	Impaired Glucose Tolerance-140 to 199. ~Diabetes>= 200.	mg/dL

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.

BILIRUBIN (TOTAL) , GEL SERUM			
BILIRUBIN (TOTAL) (Method:Vanadate oxidation)	0.70	0.3-1.2	mg/dL
ALKALINE PHOSPHATASE (Method:IFCC standardization)	111	46-116	U/L
UREA,BLOOD (Method:Urease with GLDH)	19.3	19-49	mg/dL
THYROID PANEL (T3, T4, TSH) , GEL SERUM			
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	1.13	0.60-1.81 ng/ml	ng/ml
T4-TOTAL (THYROXINE)	9.4	3.2-12.6	µg/dL



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**DEPARTMENT OF BIOCHEMISTRY**

Test Name	Result	Bio Ref. Interval	Unit
(Method:CLIA) TSH (THYROID STIMULATING HORMONE) (Method:CLIA)	1.853	0.55-4.78	µIU/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:

- Bugallo 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.
- 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:

- 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>
- Kalra S, Agarwal S, Aggarwal R, Ranabir S. Trimester-specific thyroid-stimulating hormone: An indian perspective. *Indian J Endocr Metab* 2018;22:1-4.

URIC ACID,BLOOD (Method:Uricase/Peroxidase)	5.50	3.5-7.2	mg/dL
POTASSIUM,BLOOD (Method:ISE INDIRECT)	3.80	3.5-5.5	mEq/L
SGPT/ALT (Method:Modified IFCC)	37	7-40	U/L
CALCIUM,BLOOD (Method:Arsenazo III)	9.30	8.7-10.4	mg/dL

*** End Of Report ***



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Gender	: M	Report Date	: 08/Jun/2024 12:35PM



DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
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Dr Neepa Chowdhury
MBBS, MD(Biochemistry)
SECTION DIRECTOR AND SENIOR CONSULTANT BIOCHEMIST
Reg no. WBMC 62456



Lab No.	: MDG/08-06-2024/SR9211871	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: TANMOY GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 41 Y 9 M 27 D	Collection Date	: 08/Jun/2024 10:39AM
Gender	: M	Report Date	: 08/Jun/2024 06:17PM




DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
URIC ACID, URINE, SPOT URINE			
URIC ACID, SPOT URINE (Method:URICASE)	<u>23.00</u>	37-92 mg/dL	mg/dL
<i>ESTIMATED TWICE</i>			

Suggested follow up

Correlate clinically

*** End Of Report ***


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



Lab No.	: MDG/08-06-2024/SR9211871	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: TANMOY GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 41 Y 9 M 27 D	Collection Date	: 08/Jun/2024 09:20AM
Gender	: M	Report Date	: 08/Jun/2024 12:57PM

**DEPARTMENT OF BIOCHEMISTRY**

Test Name	Result	Bio Ref. Interval	Unit
CHLORIDE,BLOOD (Method:ISE INDIRECT)	108	99-109	mEq/L

TOTAL PROTEIN [BLOOD] ALB:GLO RATIO , .			
TOTAL PROTEIN (Method:BIURET METHOD)	7.60	5.7-8.2 g/dL	g/dL
ALBUMIN (Method:BCG Dye Binding)	4.6	3.2-4.8 g/dL	g/dL
GLOBULIN (Method:Calculated)	3.00	1.8-3.2	g/dl
AG Ratio (Method:Calculated)	1.53	1.0-2.5	

GLYCATED HAEMOGLOBIN (HBA1C) , EDTA WHOLE BLOOD			
GLYCATED HEMOGLOBIN (HBA1C)	5.5	***FOR BIOLOGICAL REFERENCE INTERVAL DETAILS , PLEASE REFER TO THE BELOW MENTIONED REMARKS/NOTE WITH ADDITIONAL CLINICAL INFORMATION ***	%
HbA1c (IFCC) (Method:HPLC)	37.0		mmol/mol

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 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 B₁₂/ 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.

PDF Attached

LIPID PROFILE , GEL SERUM			
CHOLESTEROL-TOTAL (Method:Enzymatic)	203	Desirable: < 200 mg/dL Borderline high: 200-239 mg/dL High: > or =240 mg/dL	mg/dL
TRIGLYCERIDES (Method:GPO-Trinder)	168	Normal: < 150, BorderlineHigh::150-199,	mg/dL

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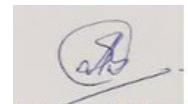


DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
HDL CHOLESTEROL (Method:Elimination/catalase)	37	High:: 200-499, VeryHigh::>500 < 40 - Low 40-59- Optimum 60 - High	mg/dl
LDL CHOLESTEROL DIRECT (Method:Elimination / Catalase)	148	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)	18	< 40 mg/dl	mg/dl
CHOL HDL Ratio (Method:Calculated)	5.5	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. Sudeshna Baral
M.B.B.S MD.
(Biochemistry)
(Consultant Biochemist)
Reg No. WBMC 64124



Lab No. : MDG/08-06-2024/SR9211871	Lab Add. : Newtown,Kolkata-700156
Patient Name : TANMOY GHOSH	Ref Dr. : Dr.MEDICAL OFFICER
Age : 41 Y 9 M 27 D	Collection Date : 08/Jun/2024 09:20AM
Gender : M	Report Date : 08/Jun/2024 02:14PM



DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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ESR (ERYTHROCYTE SEDIMENTATION RATE) , EDTA WHOLE BLOOD			
1stHour (Method:Westergren)	26	0.00 - 20.00 mm/hr	mm/hr

CBC WITH PLATELET (THROMBOCYTE) COUNT , EDTA WHOLE BLOOD			
HEMOGLOBIN (Method:PHOTOMETRIC)	15.6	13 - 17	g/dL
WBC (Method:DC detection method)	7.5	4 - 10	*10 ³ /μL
RBC (Method:DC detection method)	4.95	4.5 - 5.5	*10 ⁶ /μL
PLATELET (THROMBOCYTE) COUNT (Method:DC detection method/Microscopy)	215	150 - 450*10 ³	*10 ³ /μL
DIFFERENTIAL COUNT			
NEUTROPHILS (Method:Flowcytometry/Microscopy)	54	40 - 80 %	%
LYMPHOCYTES (Method:Flowcytometry/Microscopy)	30	20 - 40 %	%
MONOCYTES (Method:Flowcytometry/Microscopy)	10	2 - 10 %	%
EOSINOPHILS (Method:Flowcytometry/Microscopy)	05	1 - 6 %	%
BASOPHILS (Method:Flowcytometry/Microscopy)	01	0-0.9%	%
CBC SUBGROUP			
HEMATOCRIT / PCV (Method:Calculated)	46.7	40 - 50 %	%
MCV (Method:Calculated)	94.2	83 - 101 fl	fl
MCH (Method:Calculated)	31.5	27 - 32 pg	pg
MCHC (Method:Calculated)	33.4	31.5-34.5 gm/dl	gm/dl
RDW - RED CELL DISTRIBUTION WIDTH (Method:Calculated)	13.8	11.6-14%	%
PDW-PLATELET DISTRIBUTION WIDTH (Method:Calculated)	18.1	8.3 - 25 fL	fL
MPV-MEAN PLATELET VOLUME (Method:Calculated)	10.4	7.5 - 11.5 fl	fl

BLOOD GROUP ABO+RH [GEL METHOD] , EDTA WHOLE BLOOD	
ABO (Method:Gel Card)	B
RH (Method:Gel Card)	POSITIVE

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.



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Gender	: M	Report Date	: 08/Jun/2024 02:14PM



DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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Historical records check not performed.

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

Kaushik Dey
 Dr. KAUSHIK DEY
 MD (PATHOLOGY)
 CONSULTANT PATHOLOGIST
 Reg No. WBMC 66405

Lab No. : MDG/08-06-2024/SR9211871
Patient Name : TANMOY GHOSH
Age : 41 Y 9 M 27 D
Gender : M

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Jun/2024 12:52PM



DEPARTMENT OF X-RAY

DEPARTMENT OF RADIOLOGY
X-RAY REPORT OF CHEST (PA)

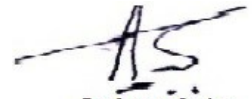
FINDINGS :

No active lung parenchymal lesion is seen.
Both the hila are normal in size, density and position.
Mediastinum is central. 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.

*** End Of Report ***


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



Lab No. : MDG/08-06-2024/SR9211871	Lab Add. : Newtown,Kolkata-700156
Patient Name : TANMOY GHOSH	Ref Dr. : Dr.MEDICAL OFFICER
Age : 41 Y 9 M 27 D	Collection Date : 08/Jun/2024 10:39AM
Gender : M	Report Date : 08/Jun/2024 03:07PM



DEPARTMENT OF CLINICAL PATHOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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Test Name	Result	Bio Ref. Interval	Unit
URINE ROUTINE ALL, ALL , URINE			
<u>PHYSICAL EXAMINATION</u>			
COLOUR	PALE YELLOW		
APPEARANCE	SLIGHTLY HAZY		
<u>CHEMICAL EXAMINATION</u>			
pH (Method:Dipstick (triple indicator method))	5.0	4.6 - 8.0	
SPECIFIC GRAVITY (Method:Dipstick (ion concentration method))	1.015	1.005 - 1.030	
PROTEIN (Method:Dipstick (protein error of pH indicators)/Manual)	NOT DETECTED	NOT DETECTED	
GLUCOSE (Method:Dipstick(glucose-oxidase-peroxidase method)/Manual)	NOT DETECTED	NOT DETECTED	
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))	NEGATIVE	NEGATIVE	
<u>MICROSCOPIC EXAMINATION</u>			
LEUKOCYTES (PUS CELLS) (Method:Microscopy)	0-1	0-5	/hpf
EPITHELIAL CELLS (Method:Microscopy)	0-1	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)	NOT DETECTED	NOT DETECTED	
YEAST (Method:Microscopy)	NOT DETECTED	NOT DETECTED	

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

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Gender	: M	Report Date	: 08/Jun/2024 03:07PM



DEPARTMENT OF CLINICAL PATHOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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and/or yeast in the urine.

*** End Of Report ***

Kaushik Dey
 Dr. KAUSHIK DEY
 MD (PATHOLOGY)
 CONSULTANT PATHOLOGIST
 Reg No. WBMC 66405

Lab No. : MDG/08-06-2024/SR9211871
Patient Name : TANMOY GHOSH
Age : 41 Y 9 M 27 D
Gender : M

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Jun/2024 03:03PM



DEPARTMENT OF CARDIOLOGY

REPORT OF E.C.G.

DATA
HEART RATE 77 Bpm
PR INTERVAL 142 Ms
QRS DURATION 100 Ms
QT INTERVAL 378 Ms
QTC INTERVAL 428 Ms

AXIS
P WAVE 48 Degree
QRS WAVE 32 Degree
T WAVE 18 Degree

IMPRESSION : Normal sinus rhythm, within normal limits.

*** End Of Report ***

Dr. A C RAY
Department of Non-invasive
Cardiology

Lab No. : MDG/08-06-2024/SR9211871
Patient Name : TANMOY GHOSH
Age : 41 Y 9 M 27 D
Gender : M

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Jun/2024 03:15PM



DEPARTMENT OF ULTRASONOGRAPHY

DEPARTMENT OF ULTRASONOGRAPHY
REPORT ON EXAMINATION OF WHOLE ABDOMEN

H/O LIVER ABSCESS DRAINING

LIVER

Liver is enlarged in size (17.15 cm) with grade II fatty changes. A hypoechoic heterogenous round shaped focal area, measuring (3.24 cmx 2.92 cm) is seen in right lobe of liver (? HEALED LIVER ABSCESS). Intrahepatic biliary radicles are not dilated. Branches of portal vein are normal

PORTA

The appearance of porta is normal. Common Bile duct is (4.8 mm) with no intraluminal pathology (Calculi /mass) could be detected at its visualised part. Portal vein is normal (7.2 mm.) at porta.

GALL BLADDER

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

PANCREAS

Echogenecity appears within limits, without any focal lesion. Shape, size & position appears normal. No Calcular disease noted. Pancreatic duct is not dilated. No peri-pancreatic collection of fluid noted.

SPLEEN

Spleen is normal in size (9.9 cm). Homogenous and smooth echotexture without any focal lesion. Splenic vein at hilum appears normal. No definite collaterals could be detected.

KIDNEYS

Both the kidneys are normal in shape, size (Rt. kidney 11.20 cm. & Lt. kidney 10.5 cm.) axes & position. Cortical echogenecity appears normal maintaining cortico-medullary & cortico-hepatic differentiation. Margin is regular and cortical thickness is uniform. No calcular disease noted. No hydronephrotic changes detected. 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. **Post void residual urine amount is 72 cc.(Significant)**

PROSTATE

Prostate is enlarged in size.Echotexture appears within normal limits. No focal alteration of its echogenecity could bedetectable.

It measures : 3.9 cm x 3.4 cm x 4.6 cm.

Approximate weight could be around = 32.9 gms

RETROPERITONEUM & PERITONEUM

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

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Report Date : 08/Jun/2024 03:15PM



DEPARTMENT OF ULTRASONOGRAPHY

IMPRESSION

- Hepatomegaly with grade II fatty liver.
- Post healed abscess.
- Prostatomegaly.
- Significant post void residual urine.

Kindly note

➤ 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.

Patient Identity not verified.

Dr. R. M. Saha

MBBS (Cal), DMCW (Cal)

PG Diploma in Ultrasonography

(Annamalai University, Tamilnadu)

REG: 43524 of WBMC

Patient Data

Sample ID: D02135770358
 Patient ID: SR9211871
 Name: TANMOY GHOSH
 Physician:
 Sex: M
 DOB:

Analysis Data

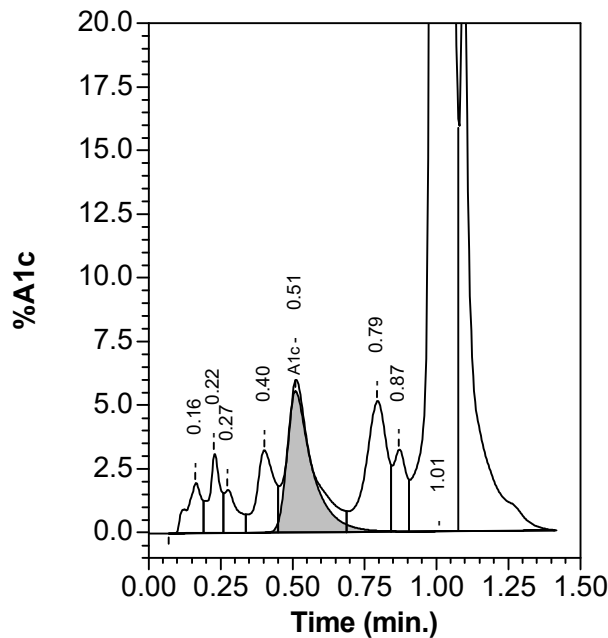
Analysis Performed: 06/08/2024 13:41:15
 Injection Number: 8455
 Run Number: 94
 Rack ID: 0007
 Tube Number: 9
 Report Generated: 06/08/2024 13:48:13
 Operator ID: ASIT

Comments:

Peak Name	NGSP %	Area %	Retention Time (min)	Peak Area
A1a	---	0.9	0.160	20829
A1b	---	1.1	0.225	24726
F	---	0.7	0.273	16101
LA1c	---	1.9	0.401	42738
A1c	5.5	---	0.510	104011
P3	---	3.5	0.792	80254
P4	---	1.3	0.867	30009
Ao	---	86.0	1.008	1956838

Total Area: 2,275,507

HbA1c (NGSP) = 5.5 % HbA1c (IFCC) = 37 mmol/mol





Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	: 08/Jun/2024 09:32AM
Gender	: F	Report Date	: 08/Jun/2024 12:15PM

**DEPARTMENT OF BIOCHEMISTRY**

Test Name	Result	Bio Ref. Interval	Unit
CREATININE, BLOOD , GEL SERUM (Method:Jaffe, alkaline picrate, kinetic)	0.66	0.5-1.1	mg/dL
GLUCOSE,FASTING (Method:Gluc Oxidase Trinder)	86	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.

PHOSPHORUS-INORGANIC,BLOOD (Method:Phosphomolybdate/UV)	2.8	2.4-5.1 mg/dL	mg/dL
THYROID PANEL (T3, T4, TSH) , GEL SERUM			
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	0.83	0.60-1.81 ng/ml	ng/ml
T4-TOTAL (THYROXINE) (Method:CLIA)	6.8	3.2-12.6	µg/dL
TSH (THYROID STIMULATING HORMONE) (Method:CLIA)	3.931	0.55-4.78	µIU/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:

- 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.
- 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:

- 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>
- Kalra S, Agarwal S, Aggarwal R, Ranabir S. Trimester-specific thyroid-stimulating hormone: An indian perspective. *Indian J Endocr Metab* 2018;22:1-4.

UREA,BLOOD (Method:Urease with GLDH)	19.3	19-49	mg/dL
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Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	: 08/Jun/2024 09:32AM
Gender	: F	Report Date	: 08/Jun/2024 12:15PM



DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
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GLUCOSE,PP (Method:Gluc Oxidase Trinder)	95	Impaired Glucose Tolerance-140 to 199.~Diabetes>= 200.	mg/dL
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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.

SODIUM,BLOOD (Method:ISE INDIRECT)	139	132 - 146	mEq/L
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CALCIUM,BLOOD (Method:Arsenazo III)	9.40	8.7-10.4	mg/dL
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URIC ACID,BLOOD (Method:Uricase/Peroxidase)	3.90	2.6-6.0	mg/dL
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POTASSIUM,BLOOD (Method:ISE INDIRECT)	3.90	3.5-5.5	mEq/L
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*** End Of Report ***

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



Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	: 08/Jun/2024 09:32AM
Gender	: F	Report Date	: 08/Jun/2024 12:35PM

**DEPARTMENT OF BIOCHEMISTRY**

Test Name	Result	Bio Ref. Interval	Unit
CHLORIDE,BLOOD (Method:ISE INDIRECT)	108	99-109	mEq/L

TOTAL PROTEIN [BLOOD] ALB:GLO RATIO , .			
TOTAL PROTEIN (Method:BIURET METHOD)	8.20	5.7-8.2 g/dL	g/dL
ALBUMIN (Method:BCG Dye Binding)	4.6	3.2-4.8 g/dL	g/dL
GLOBULIN (Method:Calculated)	3.60	1.8-3.2	g/dl
AG Ratio (Method:Calculated)	1.28	1.0-2.5	

GLYCATED HAEMOGLOBIN (HBA1C) , EDTA WHOLE BLOOD			
GLYCATED HEMOGLOBIN (HBA1C)	5.7	***FOR BIOLOGICAL REFERENCE INTERVAL DETAILS , PLEASE REFER TO THE BELOW MENTIONED REMARKS/NOTE WITH ADDITIONAL CLINICAL INFORMATION ***	%
HbA1c (IFCC) (Method:HPLC)	38.0		mmol/mol

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 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 B₁₂/ 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.

PDF Attached

LIPID PROFILE , GEL SERUM			
CHOLESTEROL-TOTAL (Method:Enzymatic)	134	Desirable: < 200 mg/dL Borderline high: 200-239 mg/dL High: > or =240 mg/dL	mg/dL
TRIGLYCERIDES (Method:GPO-Trinder)	85	Normal: < 150, BorderlineHigh::150-199,	mg/dL

Lab No. : MDG/08-06-2024/SR9211982

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Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	: 08/Jun/2024 09:32AM
Gender	: F	Report Date	: 08/Jun/2024 12:35PM

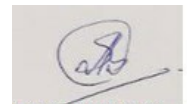


DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
HDL CHOLESTEROL (Method:Elimination/catalase)	33	High:: 200-499, VeryHigh::>500 < 40 - Low 40-59- Optimum 60 - High	mg/dl
LDL CHOLESTEROL DIRECT (Method:Elimination / Catalase)	86	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)	15	< 40 mg/dl	mg/dl
CHOL HDL Ratio (Method:Calculated)	4.1	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. Sudeshna Baral
M.B.B.S MD.
(Biochemistry)
(Consultant Biochemist)
Reg No. WBMC 64124



Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	: 08/Jun/2024 09:32AM
Gender	: F	Report Date	: 08/Jun/2024 02:14PM

**DEPARTMENT OF HAEMATOLOGY**

Test Name	Result	Bio Ref. Interval	Unit
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ESR (ERYTHROCYTE SEDIMENTATION RATE) , EDTA WHOLE BLOOD			
1stHour (Method:Westergren)	32	0.00 - 20.00 mm/hr	mm/hr

CBC WITH PLATELET (THROMBOCYTE) COUNT , EDTA WHOLE BLOOD			
HEMOGLOBIN (Method:PHOTOMETRIC)	9.0	12 - 15	g/dL
WBC (Method:DC detection method)	3.2	4 - 10	*10 ³ /μL
RBC (Method:DC detection method)	4.05	3.8 - 4.8	*10 ⁶ /μL
PLATELET (THROMBOCYTE) COUNT (Method:DC detection method/Microscopy)	179	150 - 450*10 ³	*10 ³ /μL
<u>DIFFERENTIAL COUNT</u>			
NEUTROPHILS (Method:Flowcytometry/Microscopy)	41	40 - 80 %	%
LYMPHOCYTES (Method:Flowcytometry/Microscopy)	48	20 - 40 %	%
MONOCYTES (Method:Flowcytometry/Microscopy)	09	2 - 10 %	%
EOSINOPHILS (Method:Flowcytometry/Microscopy)	02	1 - 6 %	%
BASOPHILS (Method:Flowcytometry/Microscopy)	00	0-0.9%	%
<u>CBC SUBGROUP</u>			
HEMATOCRIT / PCV (Method:Calculated)	28.2	36 - 46 %	%
MCV (Method:Calculated)	69.7	83 - 101 fl	fl
MCH (Method:Calculated)	22.1	27 - 32 pg	pg
MCHC (Method:Calculated)	31.8	31.5-34.5 gm/dl	gm/dl
RDW - RED CELL DISTRIBUTION WIDTH (Method:Calculated)	18.4	11.6-14%	%
PDW-PLATELET DISTRIBUTION WIDTH (Method:Calculated)	16.2	8.3 - 25 fL	fL
MPV-MEAN PLATELET VOLUME (Method:Calculated)	8.8	7.5 - 11.5 fl	

BLOOD GROUP ABO+RH [GEL METHOD] , EDTA WHOLE BLOOD	
ABO (Method:Gel Card)	AB
RH (Method:Gel Card)	POSITIVE

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.



Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
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Gender	: F	Report Date	: 08/Jun/2024 02:14PM



DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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Historical records check not performed.

*** End Of Report ***

Kaushik Dey
 Dr. KAUSHIK DEY
 MD (PATHOLOGY)
 CONSULTANT PATHOLOGIST
 Reg No. WBMC 66405

Lab No. : MDG/08-06-2024/SR9211982
Patient Name : SHILPI GHOSH
Age : 37 Y 11 M 21 D
Gender : F

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Jun/2024 12:51PM



DEPARTMENT OF X-RAY

DEPARTMENT OF RADIOLOGY
X-RAY REPORT OF RT KNEE (AP/LAT)

FINDINGS:


No bony injury / abnormality is seen.

No lytic / sclerotic lesion seen

Joint spaces diminished medially

Soft tissues appear normal.

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


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



Lab No. : MDG/08-06-2024/SR9211982	Lab Add. : Newtown,Kolkata-700156
Patient Name : SHILPI GHOSH	Ref Dr. : Dr.MEDICAL OFFICER
Age : 37 Y 11 M 21 D	Collection Date : 08/Jun/2024 10:30AM
Gender : F	Report Date : 08/Jun/2024 03:08PM



DEPARTMENT OF CLINICAL PATHOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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URINE ROUTINE ALL, ALL , URINE

PHYSICAL EXAMINATION

COLOUR PALE YELLOW
 APPEARANCE HAZY

CHEMICAL EXAMINATION

pH (Method:Dipstick (triple indicator method))	5.0	4.6 - 8.0	
SPECIFIC GRAVITY (Method:Dipstick (ion concentration method))	1.005	1.005 - 1.030	
PROTEIN (Method:Dipstick (protein error of pH indicators)/Manual)	NOT DETECTED	NOT DETECTED	
GLUCOSE (Method:Dipstick(glucose-oxidase-peroxidase method)/Manual)	NOT DETECTED	NOT DETECTED	
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))	POSITIVE(+)	NEGATIVE	

MICROSCOPIC EXAMINATION

LEUKOCYTES (PUS CELLS) (Method:Microscopy)	18-20	0-5	/hpf
EPITHELIAL CELLS (Method:Microscopy)	4-6	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)	PRESENT(+)	NOT DETECTED	
YEAST (Method:Microscopy)	NOT DETECTED	NOT DETECTED	

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

Lab No. : MDG/08-06-2024/SR9211982



Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	: Newtown,Kolkata-700156
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	: 08/Jun/2024 10:30AM
Gender	: F	Report Date	: 08/Jun/2024 03:08PM



DEPARTMENT OF CLINICAL PATHOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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and/or yeast in the urine.

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

Kaushik Dey
 Dr. KAUSHIK DEY
 MD (PATHOLOGY)
 CONSULTANT PATHOLOGIST
 Reg No. WBMC 66405

Lab No. : MDG/08-06-2024/SR9211982
Patient Name : SHILPI GHOSH
Age : 37 Y 11 M 21 D
Gender : F

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Jun/2024 03:02PM



DEPARTMENT OF CARDIOLOGY

REPORT OF E.C.G.

DATA
HEART RATE 59 Bpm
PR INTERVAL 144 Ms
QRS DURATION 72 Ms
QT INTERVAL 426 Ms
QTC INTERVAL 422 Ms

AXIS
P WAVE 71 Degree
QRS WAVE 77 Degree
T WAVE 26 Degree

IMPRESSION : Normal sinus rhythm, within normal limits.

*** End Of Report ***

Dr. A C RAY
Department of Non-invasive
Cardiology

Lab No.	: MDG/08-06-2024/SR9211982	Lab Add.	:
Patient Name	: SHILPI GHOSH	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 37 Y 11 M 21 D	Collection Date	:
Gender	: F	Report Date	: 08/Jun/2024 03:06PM



DEPARTMENT OF ULTRASONOGRAPHY

DEPARTMENT OF ULTRASONOGRAPHY
REPORT ON EXAMINATION OF WHOLE ABDOMEN

LIVER

Liver is normal in size (10.89 cm) having normal shape, regular smooth outline and of homogeneous echotexture. 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 (3.2 mm) with no intraluminal pathology (Calculi /mass) could be detected at its visualised part. Portal vein is normal (7.0 mm) at porta.

GALL BLADDER

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

PANCREAS

Echogenicity appears within limits, without any focal lesion. Shape, size & position appears normal. No Calcular disease noted. Pancreatic duct is not dilated. No peri-pancreatic collection of fluid noted.

SPLEEN

Spleen is normal in size (9.54 cm). Homogenous and smooth echotexture without any focal lesion. Splenic vein at hilum appears normal. No definite collaterals could be detected.

KIDNEYS

Both the kidneys are normal in shape, size (Rt. kidney 10.37 cm. & Lt. kidney 9.68 cm.) axes & position. Cortical echogenicity appears normal maintaining cortico-medullary & cortico-hepatic differentiation. Margin is regular and cortical thickness is uniform. No calcular disease noted. No hydronephrotic changes detected. 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.

UTERUS

Uterus is anteverted, normal in size (7.41 cm x 4.73 cm x 6.51 cm.) Endometrium (8.7 mm) is in midline. Myometrium appears smooth & homogenous without any detectable/sizable focal lesion.

Cervix looks normal.

Pouch of Douglas is free.

ADNEXA

Adnexa appear clear with no obvious mass lesion could be detected.

OVARIES

Right ovary is not visualised.

Left ovary is normal in size, shape, position, margin and echotexture.

Left Ovary measures 2.6 cm x 2.4 cm.

RETROPERITONEUM & PERITONEUM

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

Lab No. : MDG/08-06-2024/SR9211982

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Lab No. : MDG/08-06-2024/SR9211982
Patient Name : SHILPI GHOSH
Age : 37 Y 11 M 21 D
Gender : F

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Jun/2024 03:06PM



DEPARTMENT OF ULTRASONOGRAPHY

IMPRESSION

Sonographic study of whole abdomen does not reveal any significant abnormality.

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.

Patient Identity not verified.

Dr. R. M. Saha

MBBS (Cal), DMCW (Cal)

PG Diploma in Ultrasonography

(Annamalai University, Tamilnadu)

REG: 43524 of WBMC

Patient Data

Sample ID: D02135770385
 Patient ID: SR9211982
 Name: SHILPI GHOSH
 Physician:
 Sex: F
 DOB:

Analysis Data

Analysis Performed: 06/08/2024 13:42:52
 Injection Number: 8456
 Run Number: 94
 Rack ID: 0007
 Tube Number: 10
 Report Generated: 06/08/2024 13:48:24
 Operator ID: ASIT

Comments:

Peak Name	NGSP %	Area %	Retention Time (min)	Peak Area
A1a	---	1.3	0.165	24677
A1b	---	0.8	0.227	15243
F	---	1.4	0.276	25994
LA1c	---	1.9	0.403	33940
A1c	5.7	---	0.515	84354
P3	---	3.5	0.793	63800
P4	---	1.4	0.869	25065
Ao	---	85.1	1.011	1558901

Total Area: 1,831,975

HbA1c (NGSP) = 5.7 % HbA1c (IFCC) = 38 mmol/mol

