



Lab No.	: SG2/07-11-2024/SR9872347	Lab Add.	: Sevoke Road, Siliguri 734001
Patient Name	: MALLIKA SAHA MALLICK	Ref Dr.	: Dr.MEDICAL OFFICER
Age	: 41 Y 3 M 3 D	Collection Date	: 07/Nov/2024 11:27AM
Gender	: F	Report Date	: 07/Nov/2024 04:57PM



DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
GLUCOSE,FASTING , BLOOD, NAF PLASMA (Method:HEXOKINASE)	93	70 - 100	mg/dL
PHOSPHORUS-INORGANIC,BLOOD (Method:UV PHOSPHOMOLYBDATE)	3.4	2.5 - 4.5	mg/dL
*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)	36		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 D 10
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 glycosylated hemoglobin measurement: the position of the IFCC Working Group. Clin Chem Lab Med. 2007;45(8):1077-1080.

[PDF Attached](#)

CALCIUM,BLOOD (Method:OCPC)	8.81	8.6-10.0 mg/dl	mg/L
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*THYROID PANEL (T3, T4, TSH) , GEL SERUM			
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	0.75	0.60 - 1.81	ng/ml
T4-TOTAL (THYROXINE) (Method:CLIA)	6.9	4.5 - 10.9	microgram/dl
TSH (THYROID STIMULATING HORMONE)	2.62	0.35 - 5.5	µIU/mL



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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
(Method:CLIA)			

BIOLOGICAL REFERENCE INTERVAL : [ONLY FOR PREGNANT MOTHERS]

Trimester specific TSH LEVELS during pregnancy:

FIRST TRIMESTER	: 0.10 - 2.50 μ IU/mL
SECOND TRIMESTER	: 0.20 - 3.00 μ IU/mL
THIRD TRIMESTER	: 0.30 - 3.00 μ IU/mL

References :

1. Indian Thyroid Society guidelines for management of thyroid dysfunction during pregnancy. Clinical Practice Guidelines, New Delhi: Elsevier; 2012.
2. Stagnaro-Green A, Abalovich M, Alexander E, Azizi F, Mestman J, Negro R, et al. Guidelines of the American Thyroid Association for the Diagnosis and Management of Thyroid Disease During Pregnancy and Postpartum. Thyroid 2011;21:1081-25.
3. Dave A, Maru L, Tripathi M. Importance of Universal screening for thyroid disorders in first trimester of pregnancy. Indian J Endocr Metab [serial online] 2014 [cited 2014 Sep 25];18:735-8. Available from: <http://www.ijem.in/text.asp?2014/18/5/735/139221>.

ALKALINE PHOSPHATASE (Method:P-NPP,AMP BUFFER)	73	46 - 116	U/L
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CHLORIDE,BLOOD (Method:ISE INDIRECT)	106	98 - 107	mEq/L
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GLUCOSE,PP (Method:Hexokinase Method)	116	75-140	mg/dl
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SGOT/AST (Method:UV WITH P5P)	21	15 - 37	U/L
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SODIUM,BLOOD (Method:ISE INDIRECT)	135	136 - 145	mEq/L
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*TOTAL PROTEIN [BLOOD] ALB:GLO RATIO , .			
TOTAL PROTEIN (Method:BIURET METHOD)	7.41	6.6 - 8.7	g/dL
ALBUMIN (Method:BCP)	4	3.4-5.0 g/dl	g/dl
GLOBULIN (Method:Calculated)	3.43	1.8-3.2	g/dl
AG Ratio (Method:Calculated)	1.16	1.0 - 2.5	

*BILIRUBIN (TOTAL) , GEL SERUM			
BILIRUBIN (TOTAL) (Method:DIAZONIUM ION)	0.48	0.2 - 1.2	mg/dL

LIPID PROFILE , GEL SERUM			
CHOLESTEROL-TOTAL (Method:CHOLESTEROL OXIDASE, ESTERASE,PEROXIDASE)	203	Desirable: < 200 mg/dL Borderline high: 200-239 High: > or =240 mg/dL	mg/dL
TRIGLYCERIDES (Method:ENZYMATIC, END POINT)	135	NORMAL < 150 BORDERLINE HIGH 150-199 HIGH 200-499 VERY HIGH > 500	mg/dL
HDL CHOLESTEROL (Method:DIRECT MEASURE-PEG)	61	NO RISK : >60 mg/dL, MODERATE RISK : 40-60 mg/dL, HIGH RISK : <40	mg/dL

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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
LDL CHOLESTEROL DIRECT (Method:DIRECT MEASURE)	127	mg/dL 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)	14	< 40	mg/dL
CHOL HDL Ratio (Method:Calculated)	3.3	LOW RISK 3.3-4.4 AVERAGE RISK 4.47-7.1 MODERATE RISK 7.1-11.0 HIGH RISK >11.0	
BILIRUBIN (DIRECT) (Method:DIAZOTIZATION)	0.07	< 0.2	mg/dL
SGPT/ALT (Method:UV WITH P5P)	22	16- 63	U/L
POTASSIUM,BLOOD (Method:ISE INDIRECT)	4.82	3.5 - 5.1	mEq/L
UREA,BLOOD (Method:UREASE-COLORIMETRIC)	17	12.8 - 42.8	mg/dl
CREATININE, BLOOD (Method: ALKALINE PICRATE)	0.56	0.5 - 1.1	mg/L
URIC ACID,BLOOD (Method:URICASE ,COLORIMETRIC)	5.06	2.6 - 6.0	mg/dL

*** End Of Report ***

Dr. Ankush Chakraborty
MBBS, MD (Path), IFCAP
Consultant Pathologist
Reg. No. 65992 (WRMC)



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Age	: 41 Y 3 M 3 D	Collection Date	: 07/Nov/2024 11:27AM
Gender	: F	Report Date	: 07/Nov/2024 07:46PM



DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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BLOOD GROUP ABO+RH [GEL METHOD] , EDTA WHOLE BLOOD			
ABO (Method:Gel Card)	O		
RH (Method:Gel Card)	POSITIVE		

Gel technology Dia Med ID Micro typing system is the latest technology in transfusion Medicine. It gives more reproducible and standardized test results. It more repaid, reliable, very sensitive and objective , and hence more consistent and comparable results are obtained. Single used cards are individualised for every patient and results can be photographed / scanned and stored for future use. Special instruments that are used only for this technology also reduce risk of any contamination.

Ref:- WHO technical manual on transfusion medicine-Second Edition 2003

(RESULTS ALSO VERIFIED BY : FORWARD AND REVERSE GROUPING (TUBE AND SLIDE METHOD))

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.

CBC WITH PLATELET (THROMBOCYTE) COUNT , EDTA WHOLE BLOOD			
HEMOGLOBIN (Method:SLS haemoglobin method)	10.2	12 - 15	g/dL
WBC (Method:DC detection method)	4.6	4 - 10	*10 ³ /μL
RBC (Method:DC detection method)	4.06	3.8 - 4.8	*10 ⁶ /μL
PLATELET (THROMBOCYTE) COUNT (Method:DC detection method/Microscopy)	224	150 - 450*10 ³	*10 ³ /μL
DIFFERENTIAL COUNT			
NEUTROPHILS (Method:Flowcytometry/Microscopy)	40	40 - 80	%
LYMPHOCYTES (Method:Flowcytometry/Microscopy)	56	20 - 40	%
MONOCYTES (Method:Flowcytometry/Microscopy)	02	2 - 10	%
EOSINOPHILS (Method:Flowcytometry/Microscopy)	02	1 - 6	%
BASOPHILS (Method:Flowcytometry/Microscopy)	00	0-0.9	%
CBC SUBGROUP			
HEMATOCRIT / PCV (Method:Calculated)	32.9	36 - 46 %	%
MCV (Method:Calculated)	80.9	83 - 101 fl	fl
MCH (Method:Calculated)	25.1	27 - 32 pg	pg
MCHC (Method:Calculated)	31	31.5-34.5 gm/dl	gm/dl

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DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit
RDW - RED CELL DISTRIBUTION WIDTH (Method:Calculated)	16.3	11.6-14%	%
PDW-PLATELET DISTRIBUTION WIDTH (Method:Calculated)	23.2	8.3 - 25 fL	fL
MPV-MEAN PLATELET VOLUME (Method:Calculated)	13.0	7.5 - 11.5 fl	
RBC	NORMOCYTIC HYPOCHROMIC, MILD ANISOPOIKILOCYTOSIS.		
WBC.	RELATIVE LYMPHOCYTOSIS.		
PLATELET	ADEQUATE ON SMEAR.		

ESR (ERYTHROCYTE SEDIMENTATION RATE) , EDTA WHOLE BLOOD			
1stHour (Method:Westergren)	22	0.00 - 20.00 mm/hr	mm/hr

*** End Of Report ***

Dr. Ankush Chakraborty
MBBS, MD (Path), IFCAP
Consultant Pathologist
Reg. No. 65992 (WBMC)

Lab No. : SG2/07-11-2024/SR9872347
Patient Name : MALLIKA SAHA MALLICK
Age : 41 Y 3 M 3 D
Gender : F

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 07/Nov/2024 02:10PM



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 Sastri
MBBS, DMRT(CAL)
CONSULTANT RADIOLOGIST
Registration No.: WB-36628



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Patient Name : MALLIKA SAHA MALLICK	Ref Dr. : Dr.MEDICAL OFFICER
Age : 41 Y 3 M 3 D	Collection Date : 07/Nov/2024 02:50PM
Gender : F	Report Date : 07/Nov/2024 06:21PM



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 SLIGHTLY HAZY

CHEMICAL EXAMINATION

pH (Method:Dipstick (triple indicator method))	6.0	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)	ABSENT	NOT DETECTED
GLUCOSE (Method:Dipstick(glucose-oxidase-peroxidase method)/Manual)	ABSENT	NOT DETECTED
KETONES (ACETOACETIC ACID, ACETONE) (Method:Dipstick (Legals test)/Manual)	ABSENT	NOT DETECTED
BLOOD (Method:Dipstick (pseudoperoxidase reaction))	NEGATIVE	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

MICROSCOPIC EXAMINATION

LEUKOCYTES (PUS CELLS) (Method:Microscopy)	1-2	0-5	/hpf
EPITHELIAL CELLS (Method:Microscopy)	7-8	0-5	/hpf
RED BLOOD CELLS (Method:Microscopy)	ABSENT	0-2	/hpf
CAST (Method:Microscopy)	ABSENT	NOT DETECTED	
CRYSTALS (Method:Microscopy)	ABSENT	NOT DETECTED	
BACTERIA (Method:Microscopy)	FEW	NOT DETECTED	
YEAST (Method:Microscopy)	ABSENT	NOT DETECTED	
OTHERS	ABSENT		

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

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DEPARTMENT OF CLINICAL PATHOLOGY

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

*** End Of Report ***

Dr. Ankush Chakraborty
MBBS, MD (Path), IFCAP
Consultant Pathologist
Reg. No. 65992 (WRMC)

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Age : 41 Y 3 M 3 D
Gender : F

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 07/Nov/2024 02:26PM



DEPARTMENT OF CARDIOLOGY

DEPARTMENT OF CARDIOLOGY
REPORT OF E.C.G.

HEART RATE : 83 /min.
RHYTHM : Regular sinus.
P-WAVE : Normal
P - R INTERVAL : 160 ms,
QRS DURATION : 80 ms
QRS CONFIGURATION : NORMAL
QRS VOLTAGE : R/S in V1 4/4 mm.
R/S in V6 12/1 mm.
QRS AXIS : Normal
Q- Waves : No significant Q-wave.
QT TIME : Normal.
ST SEGMENT : Normal.
T WAVE : NORMAL
ROTATION : Normal.
OTHER FINDINGS : Nil.
IMPRESSION : ECG WITHIN NORMAL LIMIT.

*** End Of Report ***

ALG
Dr. A C RAY
Department of Non Invasive
Cardiology

Lab No. : SG2/07-11-2024/SR9872347
Patient Name : MALLIKA SAHA MALLICK
Age : 41 Y 3 M 3 D
Gender : F

Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Nov/2024 12:22PM



DEPARTMENT OF CARDIOLOGY

DEPARTMENT OF RESPIRATORY MEDICINE
REPORT OF PULMONARY FUNCTION TEST

	PRE					
	Pred	Best	% Pred	Meas 1	Meas 2	Meas 3
FVC	2.73	3.49	128	3.49	2.83	2.40
FEV 1.0	2.34	2.73	117	2.73	2.41	2.30
FEV1.0/FVC	81	78	96	78	85	96
FEF25-75%	3.42	2.44	71	2.44	2.53	3.10
PEF	5.97	5.05	85	5.05	5.03	5.28
MEF 75%	5.43	4.97	92	4.92	4.76	5.09
MEF 50%	3.83	2.98	78	2.98	3.08	3.66
MEF 25%	1.67	1.14	68	1.14	1.20	1.62

IMPRESSION :
NORMAL PULMONARY FUNCTION.

*** End Of Report ***


Dr. ARABINDA SAHA (MD,DM)
CONSULTANT CARDIOLOGIST

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Lab Add. :
Ref Dr. : Dr.MEDICAL OFFICER
Collection Date :
Report Date : 08/Nov/2024 12:20PM



DEPARTMENT OF CARDIOLOGY

DEPARTMENT OF CARDIOLOGY
REPORT ON EXAMINATION OF STRESS TEST (T.M.T)

RESULT : FAIR EXERCISE(7.0METS) TOLERANCE. NORMAL BP &
HEART RATE RESPONSE. NO SIGNIFICANT ST-T
SEGMENT CHANGE IN LEADS.

THE TEST TERMINATED BECAUSE OF SOB.

IMPRESSION : THE TEST NEGATIVE FOR INDUCIBLE ISCHAEMIA.

Thank you for the opportunity to participate in the care of your patient.

*** End Of Report ***


Dr. ARABINDA SAHA (MD,DM)
CONSULTANT CARDIOLOGIST

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DEPARTMENT OF ULTRASONOGRAPHY

DEPARTMENT OF ULTRASONOGRAPHY
REPORT ON EXAMINATION OF WHOLE ABDOMEN

LIVER

Liver is normal in size 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 normal with no intraluminal pathology (Calculi /mass) could be detected at its visualised part. Portal vein is normal at porta.

GALL BLADDER

Gallbladder is operated.

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. 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 93 mm. & Lt. kidney 96 mm.) 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. 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 (78 mm. x 37 mm. x 40 mm). Endometrium (6 mm) is in midline. Myometrium appears smooth & homogenous without any detectable/sizable focal lesion. Cervix looks normal. Pouch of Douglas is free.

OVARIES

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

Thin walled anechoic cyst (34 x 29 mm) seen at left ovary.

Right ovary measures 31 x 20 mm.

IMPRESSION :

- i) Post cholecystectomy status.
- ii) Thin walled anechoic cyst (34 x 29 mm) at left ovary.

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DEPARTMENT OF ULTRASONOGRAPHY

(Please correlate clinically & with other investigation. Follow up suggested).

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.

*** End Of Report ***

MS
DR. MUKTI SARKAR MD.
CONSULTANT RADIOLOGIST

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Age	: 41 Y 3 M 3 D	Collection Date	:
Gender	: F	Report Date	: 08/Nov/2024 01:24PM



DEPARTMENT OF MAMMOGRAPHY

DEPARTMENT OF RADIOLOGY
MAMMOGRAPHY OF BOTH BREAST

Cranio-caudal & medio-lateral oblique views of both mammary gland are taken along with axillary tail.

Reveal coarse texture of glandular elements mixed with fatty tissue.

No macro / micro calcification noted.

Skin & nipple outline are normal on both sides.

Multiple small nodes noted at both axilla (4 to 5 mm).

IMPRESSION :

Normal mammography of both breast.

N.B: Mammography may be normal in fibroadenosis.

(Please correlate clinically & with other investigation. Follow up suggested).

Breast imaging and data system

Category 0: Need additional imaging

Category 1: Negative category 2: Benign findings

Category 3: Probably benign (< 2 % risk of malignancy)

short interval follow up suggested (in 6 months)

Category 4: Suspicious abnormality - biopsy should be considered

Category 5: Highly suggestive of malignancy

Appropriate action should be taken

Category 6: Known biopsy proven malignancy]

[INFORMATION REGARDING MAMMOGRAMS

1.A report that is negative for malignancy should not delay biopsy if there is a dominant or clinically suspicious mass.

2.In dense breasts an underlying mass lesion may be obscured.

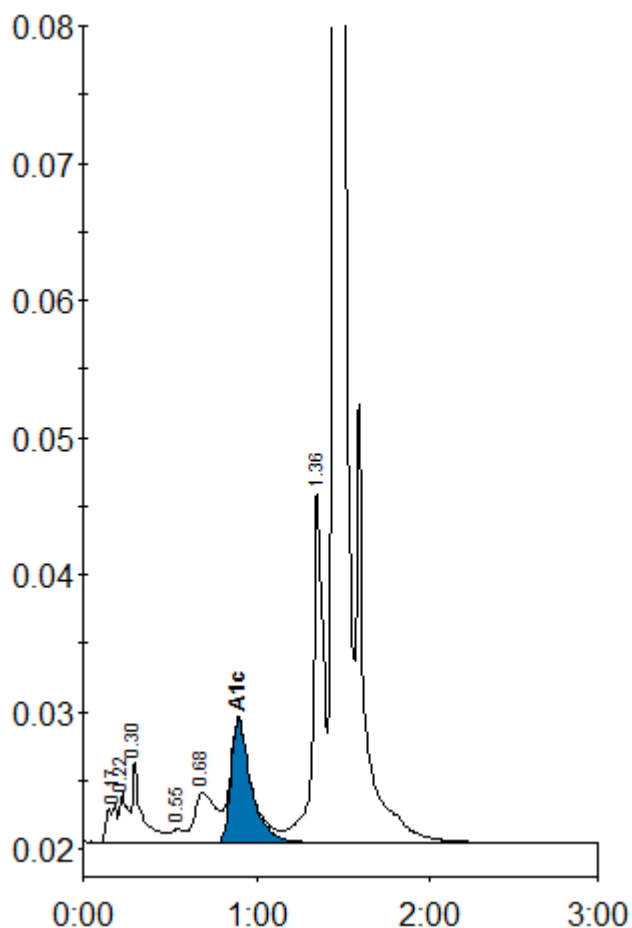
3.False positive diagnoses of cancer may occur in small percentage of case.]

MS

DR. MUKTI SARKAR MD.
CONSULTANT RADIOLOGIST

Patient report

Sample ID: E02132880682
 Injection date 07/11/2024 04:44 PM
 Injection #: 24 D-10 Method: HbA1c
 Rack #: --- Rack position: 6
 Bio-Rad v: 5.00-2 S/N: #DM23F10804



Peak table - ID: E02132880682

Peak	R.time	Height	Area	Area %
Unknown	0.17	2397	10559	0.6
A1a	0.22	3610	12085	0.6
A1b	0.30	6038	23363	1.2
F	0.55	982	5794	0.3
LA1c/CHb-1	0.68	3701	30439	1.6
A1c	0.90	9066	72992	5.5
P3	1.36	25408	107579	5.7
A0	1.45	669341	1635063	86.2
Total Area:			1897874	

Concentration:	%	mmol/mol
A1c	5.5	36