

Patient Name : SNEHA AGARWAL

Age : 35 Y 8 M 20 D

Gender

: F

Lab Add. : Sevoke Road, Siliguri 734001

: Dr.MEDICAL OFFICER

: 24/Sep/2024 11:12AM

Report Date : 24/Sep/2024 06:00PM



DEPARTMENT OF BIOCHEMISTRY

Ref Dr.

Collection Date

Test Name	Result	Bio Ref. Interval	Unit
ALKALINE PHOSPHATASE , GEL SERUM (Method:P-NPP,AMP BUFFER)	<u>119</u>	46 - 116	U/L
*BILIRUBIN (TOTAL) , GEL SERUM			
BILIRUBIN (TOTAL) (Method:DIAZONIUM ION)	0.63	0.2 - 1.2	mg/dL
SGPT/ALT (Method:UV WITH P5P)	39	16- 63	U/L
SODIUM,BLOOD (Method:ISE INDIRECT)	134	136 - 145	mEq/L
POTASSIUM,BLOOD (Method:ISE INDIRECT)	3.91	3.5 - 5.1	mEq/L
UREA,BLOOD (Method:UREASE-COLORIMETRIC)	20	12.8 - 42.8	mg/dl
GLUCOSE,FASTING (Method:HEXOKINASE)	90	70 - 100	mg/dL
PHOSPHORUS-INORGANIC,BLOOD (Method:UV PHOSPHOMOLYBDATE)	4	2.5 - 4.5	mg/dL
URIC ACID,BLOOD (Method:URICASE ,COLORICMETRIC)	5.49	2.6 - 6.0	mg/dL
*TOTAL PROTEIN [BLOOD] ALB:GLO RA	ATIO , .		
TOTAL PROTEIN (Method:BIURET METHOD)	7.23	6.6 - 8.7	g/dL
ALBUMIN (Method:BCP)	4	3.4 -5.0 g/dl	g/dl
GLOBULIN (Method:Calculated)	<u>3.28</u>	1.8-3.2	g/dl
AG Ratio (Method:Calculated)	1.2	1.0 - 2.5	
*GLYCATED HAEMOGLOBIN (HBA1C), A	EDTA WHOLE BLOOD		
GLYCATED HEMOGLOBIN (HBA1C)	5.3	***FOR BIOLOGICAL REFERENCE INTERVAL DETAILS , PLEASE REFER TO THE BELOW MENTIONED REMARKS/NOTE	%

Clinical Information and Laboratory clinical interpretation on Biological Reference Interval:

35

 $\begin{tabular}{ll} Low risk / Normal / non-diabetic & : <5.7\% (NGSP) & / < 39 mmol/mol (IFCC) \\ Pre-diabetes/High risk of Diabetes : 5.7\%- 6.4\% (NGSP) / 39 - < 48 mmol/mol (IFCC) \\ Diabetics-HbA1c level & : >/= 6.5\% (NGSP) & / > 48 mmol/mol (IFCC) \\ \end{tabular}$

Analyzer used: Bio-Rad D 10

HbA1c (IFCC)

(Method:HPLC)

mmol/mol

WITH ADDITIONAL CLINICAL

INFORMATION ***



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DEPARTMENT OF BIOCHEMISTRY

Test Name Result Bio Ref. Interval Unit

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:CHOLESTEROL OXIDASE, ESTERASE,PEROXIDASE)	182	Desirable: < 200 mg/dL Borderline mg/dL high: 200-239 High: > or =240 mg/dL
TRIGLYCERIDES (Method:ENZYMATIC, END POINT)	<u>156</u>	NORMAL < 150 BORDERLINE HIGH mg/dL 150-199 HIGH 200-499 VERY HIGH > 500
HDL CHOLESTEROL (Method:DIRECT MEASURE-PEG)	51	NO RISK : >60 mg/dL, MODERATE mg/dL RISK : 40-60 mg/dL, HIGH RISK : <40 mg/dL
LDL CHOLESTEROL DIRECT (Method:DIRECT MEASURE)	117	OPTIMAL: <100 mg/dL, Near mg/dL optimal/ above optimal: 100-129 mg/dL, Borderline high: 130-159 mg/dL, High: 160-189 mg/dL, Very high: >=190 mg/dL
VLDL (Method:Calculated)	14	< 40 mg/dL
CHOL HDL Ratio (Method:Calculated)	3.6	LOW RISK 3.3-4.4 AVERAGE RISK 4.47-7.1 MODERATE RISK 7.1-11.0 HIGH RISK >11.0

NOTE: Elevated Triglyceride value is to be interpreted in the light of previous 72 hrs dietary intake of lipids. Repeat estimation with 72 hrs fat restricted diet followed by 12 hrs fasting, suggested for better evaluation.

CALCIUM,BLOOD (Method:OCPC)	9.47	8.6-10.0 mg/dl	mg/L	

GLUCOSE,PP 110 75-140 mg/dl



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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
(Method:Hexokinase Method)			
SGOT/AST (Method:UV WITH P5P)	25	15 - 37	U/L
CREATININE, BLOOD (Method: ALKALINE PICRATE)	0.78	0.5 - 1.1	mg/L
BILIRUBIN (DIRECT) (Method:DIAZOTIZATION)	<u>0.11</u>	< 0.2	mg/dL
CHLORIDE,BLOOD (Method:ISE INDIRECT)	104	98 - 107	mEq/L
*THYROID PANEL (T3, T4, TSH), GEL SERUM	1		
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	1.01	0.60 - 1.81	ng/ml
T4-TOTAL (THYROXINE) (Method:CLIA)	8.5	4.5 - 10.9	microgram/dl
TSH (THYROID STIMULATING HORMONE) (Method:CLIA)	1.93	0.35 - 5.5	μIU/mL

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.

*** End Of Report ***

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

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: Sevoke Road, Siliguri 734001 : Dr. MEDICAL OFFICER

MC-2176

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: SNEHA AGARWAL Ref Dr.

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 : 24/Sep/2024 11:12AM

Gender : F Report Date : 24/Sep/2024 04:59PM

DEPARTMENT OF HAEMATOLOGY

Test Name Result Bio Ref. Interval Unit

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

ABO C

(Method:Gel Card)

RH POSITIVE

(Method:Gel Card)

Patient Name

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.

ESR (ERYTHROCYTE SEDIME	NTATION RATE), EDTA WH	OLE BLOOD		
1stHour	<u>36</u>	0.00 - 20.00 mm/hr	mm/hr	
(Method:Westergren)				

CBC WITH PLATELET (THROMBOCYTE) COUNT, EDTA WHOLE BLOOD						
HEMOGLOBIN	<u>11.5</u>	12 - 15	g/dL			
(Method:SLS haemoglobin method) WBC	8.6	4 - 10	*10^3/µL			
(Method:DC detection method)	0.0	1 10	10 0/μΕ			
RBC	4.18	3.8 - 4.8	*10^6/µL			
(Method:DC detection method) PLATELET (THROMBOCYTE) COUNT (Method:DC detection method/Microscopy) DIFFERENTIAL COUNT	446	150 - 450*10^3	*10^3/µL			
NEUTROPHILS (Method:Flowcytometry/Microscopy)	60	40 - 80	%			
LYMPHOCYTES (Method:Flowcytometry/Microscopy)	36	20 - 40	%			
MONOCYTES (Method:Flowcytometry/Microscopy)	02	2 - 10	%			
EOSINOPHILS (Method:Flowcytometry/Microscopy)	02	1 - 6	%			
BASOPHILS (Method:Flowcytometry/Microscopy) CBC SUBGROUP	00	0-0.9	%			
HEMATOCRIT / PCV (Method:Calculated)	<u>34.4</u>	36 - 46 %	%			
MCV	<u>82.1</u>	83 - 101 fl	fl			

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Lab No.



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DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit	
(Method:Calculated)				
MCH	27.5	27 - 32 pg	pg	
(Method:Calculated) MCHC (Method:Calculated)	33.5	31.5-34.5 gm/dl	gm/dl	
RDW - RED CELL DISTRIBUTION WIDTH (Method:Calculated)	14.3	11.6-14%	%	
PDW-PLATELET DISTRIBUTION WIDTH (Method:Calculated)	18.6	8.3 - 25 fL	fL	
MPV-MEAN PLATELET VOLUME (Method:Calculated)	11.5	7.5 - 11.5 fl		
RBC	NORMOCYTIC NORMOCHROMIC.			
WBC.	NORMAL MORPHOLOGY.			
PLATELET	ADEQUATE ON SMEAR.			

*** End Of Report ***

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



Patient Name : SNEHA AGARWAL Ref Dr. : Dr.MEDICAL OFFICER

Age : 35 Y 8 M 20 D Collection Date

Gender : F Report Date : 24/Sep/2024 12:40PM



DEPARTMENT OF X-RAY

Lab Add.

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

FINDINGS:

- Cardiac size appears within normal limits. Margin is well visualised and cardiac silhoutte is smoothly outlined. Shape is within normal limit.
- Lung parenchyma shows no focal lesion. No general alteration of radiographic density. Apices are clear. Bronchovascular lung markings are within normal.
- · Lateral costo-phrenic angles are clear.
- Domes of diaphragm are smoothly outlined. Position is within normal limits.

IMP	RES	SSIC	NC	:
Nor	mal	stu	ıdv	

*** End Of Report ***

DR. MUKTI SARKAR MD.
CONSULTANT RADIOLOGIST

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 Age
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 Report Date
 : 25/Sep/2024 11:46AM



DEPARTMENT OF CLINICAL PATHOLOGY

Test Name Result Bio Ref. Interval Unit

URINE ROUTINE ALL, ALL, URINE			
PHYSICAL EXAMINATION			
COLOUR	STRAW		
APPEARANCE	CLEAR		
CHEMICAL EXAMINATION			
pH (Method:Dipstick (triple indicator method))	6.0	4.6 - 8.0	
SPECIFIC GRAVITY (Method:Dipstick (ion concentration method))	1.010	1.005 - 1.030	
PROTEIN (Method:Dipstick (protein error of pH	ABSENT	NOT DETECTED	
indicators)/Manual) GLUCOSE (Method:Dipstick(glucose-oxidase-peroxidase	ABSENT	NOT DETECTED	
method)/Manual) KETONES (ACETOACETIC ACID, ACETONE)	ABSENT	NOT DETECTED	
(Method:Dipstick (Legals test)/Manual) BLOOD	ABSENT	NOT DETECTED	
(Method:Dipstick (pseudoperoxidase reaction)) BILIRUBIN (Method:Dipstick (azo-diazo reaction)/Manual)	ABSENT	NEGATIVE	
UROBILINOGEN (Method:Dipstick (diazonium ion reaction)/Manual)	ABSENT	NEGATIVE	
NITRITE (Method:Dipstick (Griess test))	ABSENT	NEGATIVE	
LEUCOCYTE ESTERASE (Method:Dipstick (ester hydrolysis reaction)) MICROSCOPIC EXAMINATION	ABSENT	NEGATIVE	
LEUKOCYTES (PUS CELLS) (Method:Microscopy)	1-2	0-5	/hpf
EPITHELIAL CELLS (Method:Microscopy)	6-7	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:

- 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





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DEPARTMENT OF CLINICAL PATHOLOGY

Test Name Result Bio Ref. Interval Unit

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 (WBMC)

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Patient Name : SNEHA AGARWAL Ref Dr. : Dr.MEDICAL OFFICER

Age : 35 Y 8 M 20 D Collection Date

 Gender
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 Report Date
 : 25/Sep/2024 12:51PM



DEPARTMENT OF CARDIOLOGY

Lab Add.

DEPARTMENT OF RESPIRATORY MEDICINE REPORT OF PULMONARY FUNCTION TEST

	I	PRE				
	Pred	Best	% Pred	Meas 1	Meas 2	Meas 3
FVC	2.98	4.33	146	4.33	3.18	2.87
FEV 1.0	2.57	3.63	141	3.63	2.59	2.51
FEV1.0/FVC	82	84	102	84	81	87
FEF25-75%	3.65	3.63	99	3.63	2.49	3.07
PEF	6.26	6.96	111	6.96	6.94	6.44
MEF 75%	5.65	5.16	91	5.16	5.89	6.10
MEF 50%	4.03	4.38	109	4.38	2.90	3.72
MEF 25%	1.84	1.74	95	1.74	1.06	1.38

IMPRESSION:

NORMAL PULMONARY FUNCTION.

*** End Of Report ***

Dr. ARABINDA SAHA (MD,DM) CONSULTANT CARDIOLOGIST



Patient Name : SNEHA AGARWAL Ref Dr. : Dr.MEDICAL OFFICER

Age : 35 Y 8 M 20 D Collection Date

Gender : F Report Date : 24/Sep/2024 02:02PM

DEPARTMENT OF CARDIOLOGY

DEPARTMENT OF CARDIOLOGY REPORT OF E.C.G.

Lab Add.

HEART RATE : 73 /min.

RHYTHM : Regular sinus.

P-WAVE : Normal

P-RINTERVAL: 120 ms,

QRS DURATION : 80 ms

QRS CONFIGURATION : NORMAL

QRS VOLTAGE : R/S in V1 1/2 mm.

R/S in V6 8/2 mm.

QRS AXIS : +60°

Q- Waves : No significant Q-wave.

QCT INTERVAL : 409 ms

ST SEGMENT : Normal.

T WAVE : NORMAL

ROTATION : Normal.

OTHER FINDINGS : Nil.

IMPRESSION : ECG WITHIN NORMAL LIMIT.

*** End Of Report ***

Dr. ARABINDA SAHA (MD,DM) CONSULTANT CARDIOLOGIST

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Patient Name

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Lab Add.

Age : 35 Y 8 M 20 D Collection Date :

Gender : F Report Date : 25/Sep/2024 12:46PM



DEPARTMENT OF CARDIOLOGY

<u>DEPARTMENT OF CARDIOLOGY</u> REPORT ON EXAMINATION OF STRESS TEST (T.M.T)

RESULT: FAIR EXERCISE (10.1 METS) TOLERANCE. NORMAL

HEART RATE & BP RESPONSE. SIGNIFICANT ST-T

SEGMENT CHANGE IN LEADS II, III, avF & V4 - V6.

THE TEST TERMINATED BECAUSE OF SOB.

IMPRESSION : THE TEST <u>POSITIVE</u> 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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Lab No. : SIL/24-09-2024/SR9699679 Lab Add.

Patient Name : SNEHA AGARWAL Ref Dr. : Dr.MEDICAL OFFICER

Age : 35 Y 8 M 20 D Collection Date :

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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 **shows grade I fatty change.** 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 visualsed part. Portal vein is normal 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. 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 92 mm. & Lt. kidney 99 mm.) axes & position. Cortical echogenecity 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 (66 mm. x 30 mm. x 44 mm.) Endometrium (05mm) is in midline. Myometrium appears smooth & homogenous without any detectable/sizable focal lesion.

Cervix looks normal.Pouch of Douglas is free.

OVARIES

Ovaries are normal in size, shape, position, margin and echotexture.

Right ovary measures 23 mm x 17 mm.

Left Ovary measures 22 mm x 16 mm.

IMPRESSION:

Grade I fatty change in liver.

Kindly note

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: F Report Date : 24/Sep/2024 05:04PM Gender



- DEPARTMENT OF ULTRASONOGRAPHY > 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 ***

DR. MUKTI SARKAR MD. CONSULTANT RADIOLOGIST

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DEPARTMENT OF MAMMOGRAPHY

<u>DEPARTMENT OF RADIOLOGY</u> MAMMOGRAPHY OF BOTH BREASTS

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.

Bilateral dense breast.....no obvious focal lesion.

Skin & nipple outline are normal on both sides.

AXILLA: No nodes on both sides.

IMPRESSION:

Bilateral dense breast.....BI-RADS II.

(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.]

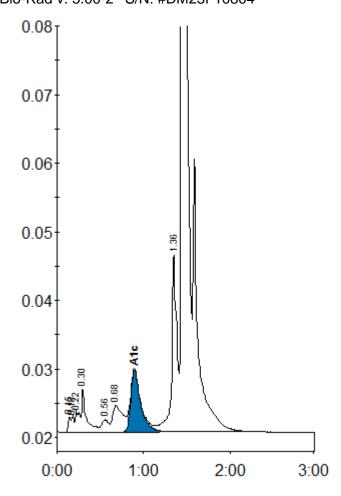
DR. MUKTI SARKAR MD.
CONSULTANT RADIOLOGIST

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Patient report

Sample ID: E02132884814

Injection date 24/09/2024 08:01 AM Injection #: 16 D-10 Method: HbA1c Rack #: --- Rack position: 2
Bio-Rad v: 5.00-2 S/N: #DM23F10804



Peak table - ID: E02132884814

Peak	R.time	Height	Area	Area %
Unknown	0.15	2158	4569	0.2
Unknown	0.18	2396	4689	0.2
A1a	0.22	3223	11447	0.6
A1b	0.30	6507	23285	1.2
F	0.56	1773	10119	0.5
LA1c/CHb-1	0.68	3845	30586	1.6
A1c	0.89	9041	73212	5.3
P3	1.36	25788	108421	5.6
A0	1.45	705086	1681120	86.3

Concentration:	%	mmol/mol
Δ1c	53	35

1947448

Total Area: