



Patient Name : CHUNKU BHUTIA

Age : 34 Y 6 M 12 D

Gender : F

Lab Add. : Sevoke Road, Siliguri 734001

Ref Dr. : Dr.MEDICAL OFFICER

Collection Date : 23/Mar/2024 09:27AM

: 23/Mar/2024 01:59PM



DEPARTMENT OF BIOCHEMISTRY

Report Date

Test Name	Result	Bio Ref. Interval	Unit
SODIUM,BLOOD , GEL SERUM (Method:ISE INDIRECT)	141	136 - 145	mEq/L
CHLORIDE,BLOOD (Method:ISE INDIRECT)	105	98 - 107	mEq/L
GLUCOSE,FASTING (Method:Hexokinase Method)	79	70 - 100	mg/dl
CALCIUM,BLOOD (Method:OCPC)	9.30	8.6-10.0 mg/dl	mg/L
*TOTAL PROTEIN [BLOOD] ALB:GLO RA	ΠΟ , .		
TOTAL PROTEIN (Method:BIURET METHOD)	7.02	6.6 - 8.7	g/dL
ALBUMIN (Method:BCP)	4.1	3.4 -5.0 g/dl	g/dl
GLOBULIN (Method:Calculated)	2.96	1.8-3.2	g/dl
AG Ratio (Method:Calculated)	1.37	1.0 - 2.5	
GLUCOSE,PP (Method:Hexokinase Method)	84	75-140	mg/dl
*THYROID PANEL (T3, T4, TSH), GEL SERUI	И		
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	0.99	0.60 - 1.81 ng/ml	ng/ml
T4-TOTAL (THYROXINE) (Method:CLIA)	8.0	4.5 - 10.9	microgram/dl
TSH (THYROID STIMULATING HORMONE) (Method:CLIA)	2.00	0.35-5.5	μlU/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.

LIPID	PROFILE.	GEL SERUM
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CHOLESTEROL-TOTAL 189 Desirable: < 200 mg/dL Borderline mg/d (Method:CHOLESTEROL OXIDASE, ESTERASE,PEROXIDASE)

Desirable: < 200 mg/dL Borderline mg/d high: 200-239 High: > or =240 mg/dL

TRIGLYCERIDES 47 NORMAL < 150 BORDERLINE HIGH mg/dl (Method:ENZYMATIC, END POINT) 150-199 HIGH 200-499 VERY HIGH >

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MC-2

 Patient Name
 : CHUNKU BHUTIA
 Ref Dr.
 : Dr.MEDICAL OFFICER

 Age
 : 34 Y 6 M 12 D
 Collection Date
 : 23/Mar/2024 09:27AM

 Gender
 : F
 Report Date
 : 23/Mar/2024 01:59PM



DEPARTMENT OF BIOCHEMISTRY

	DEFINITION OF BIOCHEMISTRY				
Test Name	Result	Bio Ref. Interval	Unit		
1		500			
HDL CHOLESTEROL	66	NO RISK : >60 mg/dL, MODERATE	ma/dl		
(Method:DIRECT MEASURE-PEG)	<u>00</u>	RISK: 40-60 mg/dL, HIGH RISK: <40			
(Mothod.Birtzo1 Mz/roortz 1 zo)		mg/dL	,		
LDL CHOLESTEROL DIRECT	109	OPTIMAL : <100 mg/dL, Near	mg/dl		
(Method:DIRECT MEASURE)	<u></u>	optimal/ above optimal: 100-129	g,		
,		mg/dL, Borderline high: 130-159			
		mg/dL, High: 160-189 mg/dL, Very			
		high:>=190 mg/dL			
VLDL	14	< 40 mg/dl	mg/dL		
(Method:Calculated)					
CHOL HDL Ratio	<u>2.9</u>	LOW RISK 3.3-4.4 AVERAGE RISK			
(Method:Calculated)		4.47-7.1 MODERATE RISK 7.1-11.0			
		HIGH RISK >11.0			
UREA,BLOOD	21.0	12.8-42.8	mg/dl		
(Method:UREASE-COLORIMETRIC)	21.0	12.0 42.0	mg/ai		
()					
URIC ACID,BLOOD	4.70	2.6 - 6.0	mg/dl		
(Method:URICASE ,COLORICMETRIC)					
*GLYCATED HAEMOGLOBIN (HBA1C)	EDTA WHOLE BLOOD				
GLYCATED HEMOGLOBIN (HBA1C)	5.3	***FOR BIOLOGICAL REFERENCE	0/_		
GETOATED HEMIOGEODIN (HBATO)	5.5	INTERVAL DETAILS , PLEASE	76		
		REFER TO THE BELOW			
		MENTIONED REMARKS/NOTE			
		WITH ADDITIONAL CLINICAL			
		INFORMATION ***			
HbA1c (IFCC)	34.0		mmol/mol		

Clinical Information and Laboratory clinical interpretation on Biological Reference Interval:

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

Analyzer used : Bio-Rad D 10 Method : HPLC Cation Exchange

(Method:HPLC)

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 Lab No. : SG2/23-03-2024/SR8904397 Page 2 of 12





: SG2/23-03-2024/SR8904397 Lab Add. : Sevoke Road, Siliguri 734001

Patient Name : CHUNKU BHUTIA Ref Dr. : Dr.MEDICAL OFFICER : 34 Y 6 M 12 D **Collection Date** : 23/Mar/2024 09:27AM Age : F Gender

: 23/Mar/2024 01:59PM Report Date



DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
			•

Diabetes. Ann Intern Med. Published online 1 March 2016. doi:10.7326/M15-3016.

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

PDF Attached

Lab No.

POTASSIUM,BLOOD (Method:ISE INDIRECT)	<u>5.20</u>	3.5 - 5.1	mEq/L	
PHOSPHORUS-INORGANIC,BLOOD (Method:UV PHOSPHOMOLYBDATE)	4.0	2.5-4.5 mg/dl	mg/dl	
CREATININE, BLOOD (Method: ALKALINE PICRATE)	0.70	0.50 - 1.10	mg/dl	

*** End Of Report ***

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

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мс-2178 : SG2/23-03-2024/SR8904397

: 34 Y 6 M 12 D

Patient Name : CHUNKU BHUTIA

Gender : F

Lab No.

Age

Lab Add. : Sevoke Road, Siliguri 734001

Ref Dr. : Dr.MEDICAL OFFICER

: 23/Mar/2024 06:45PM

Collection Date : 23/Mar/2024 09:27AM

DEPARTMENT OF HAEMATOLOGY

Report Date

Test Name Result Bio Ref. Interval Unit	
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*CBC WITH PLATELET (THROMBOCYTE)	COUNT, EDTA WHOLE BLO	OD .	
HEMOGLOBIN	13.9	12 - 15	g/dL
(Method:SLS haemoglobin method)			
WBC	5.4	4 - 10	*10^3/µL
(Method:DC detection method)			
RBC	4.22	3.8 - 4.8	*10^6/µL
(Method:DC detection method)	050	450 450*4040	*4000/ 1
PLATELET (THROMBOCYTE) COUNT	250	150 - 450*10^3	*10^3/µL
(Method:DC detection method/Microscopy) DIFFERENTIAL COUNT			
NEUTROPHILS	61	40 - 80 %	%
(Method:Flowcytometry/Microscopy)	0.5	00 40 0/	0/
LYMPHOCYTES	35	20 - 40 %	%
(Method:Flowcytometry/Microscopy) MONOCYTES	02	2 - 10 %	%
(Method:Flowcytometry/Microscopy)	02	2 - 10 %	70
EOSINOPHILS	02	1 - 6 %	%
(Method:Flowcytometry/Microscopy)	02	1 - 0 /8	76
BASOPHILS	00	0-0.9%	%
(Method:Flowcytometry/Microscopy)		0 0.070	,,
CBC SUBGROUP			
HEMATOCRIT / PCV	39.8	36 - 46 %	%
(Method:Calculated)	30.0	00 10 70	,,
MCV	94.3	83 - 101 fl	fl
(Method:Calculated)			
MCH	<u>33.0</u>	27 - 32 pg	pg
(Method:Calculated)			
MCHC	<u>34.9</u>	31.5-34.5 gm/dl	gm/dl
(Method:Calculated)			
RDW - RED CELL DISTRIBUTION WIDTH	<u>15.6</u>	11.6-14%	%
(Method:Calculated)			
PDW-PLATELET DISTRIBUTION WIDTH	18.4	8.3 - 25 fL	fL
(Method:Calculated)			
MPV-MEAN PLATELET VOLUME	11.5	7.5 - 11.5 fl	
(Method:Calculated)	NORMOCYTIC		
RBC			
MDO	NORMOCHROMIC.		
WBC.	UNREMARKABLE		
PLATELET	ADEQUATE ON SMEAR	≺	

FSR	(ERYTHROCYTE SEDIMENTATION RATE).	EDTA WHOLE BLOOD

 1stHour
 12
 0.00 - 20.00 mm/hr
 mm/hr

 (Method:Westergren)
 mm/hr
 mm/hr

*** End Of Report ***

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Lab Add. : Sevoke Road, Siliguri 734001

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Collection Date : 23/Mar/2024 09:27AM

Report Date : 23/Mar/2024 06:45PM

DEPARTMENT OF HAEMATOLOGY

Test Name Result Bio Ref. Interval Unit

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





Lab No.

: SG2/23-03-2024/SR8904397

: CHUNKU BHUTIA : 34 Y 6 M 12 D

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Lab Add. : Sevoke Road, Siliguri 734001

: Dr.MEDICAL OFFICER Ref Dr.

Collection Date : 23/Mar/2024 09:27AM

: 23/Mar/2024 05:59PM Report Date

DEPARTMENT OF HAEMATOLOGY

Test Name Result Bio Ref. Interval Unit

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

(Method:Gel Card)

RH **POSITIVE**

(Method:Gel Card)

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.

*** End Of Report ***

Dr. PRABHA ANAND MBBS, MD(Microbiology) CONSULTANT MICROBIOLOGISTS Reg No. WBMC 92308

Lab No. SG2/23-03-2024/SR8904397



Patient Name : CHUNKU BHUTIA Age

:F

: 34 Y 6 M 12 D

Lab Add.

Ref Dr.

Report Date

: Sevoke Road, Siliguri 734001

: Dr.MEDICAL OFFICER

Collection Date : 23/Mar/2024 11:13AM : 25/Mar/2024 06:22PM



DEPARTMENT OF CYTOLOGY

DEPARTMENT OF PATHOLOGY REPORT ON EXAMINATION OF CERVICAL SMEAR FOR EXFOLIATIVE CYTOLOGY

SPECIMEN TYPE:

Gender

Conventional cervical PAP smear.

SPECIMEN ADEQUACY:

Satisfactory for evaluation. Endocervical cells not seen.

GENERAL DIAGNOSTIC CATEGORIZATION:

Negative for intraepithelial lesion / malignancy [NILM].

IMPRESSION:

Reactive cellular changes associated with inflammation.

NOTE: Reported as per The 2014 Bethesda system of reporting cervical cytology.

ENCL: Two (02) slides.

*** End Of Report ***

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

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: SG2/23-03-2024/SR8904397 Lab No.



: CHUNKU BHUTIA Ref Dr. : Dr.MEDICAL OFFICER

Age : 34 Y 6 M 12 D Collection Date

Gender : F Report Date : 24/Mar/2024 11:07AM



DEPARTMENT OF X-RAY

Lab Add.

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

FINDINGS:

Patient Name

- 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.

IMPRE	ESSION	:
Norma	al study.	

*** End Of Report ***

DR. Ziaul Mustafa MD, Radiodiagnosis

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Lab No.

: SG2/23-03-2024/SR8904397 Lab Add. : Sevoke Road,Siliguri 734001

 Patient Name
 : CHUNKU BHUTIA
 Ref Dr.
 : Dr.MEDICAL OFFICER

 Age
 : 34 Y 6 M 12 D
 Collection Date
 : 23/Mar/2024 09:29AM

 Gender
 : F
 Report Date
 : 23/Mar/2024 01:34PM



DEPARTMENT OF CLINICAL PATHOLOGY

Test Name Result Bio Ref. Interval Unit

PALE YELLOW		
SLIGHTLY HAZY		
5.0	4.6 - 8.0	
1.010	1.005 - 1.030	
ADCENIT	NOT DETECTED	
ADSENT	NOT DETECTED	
ABSENT	NOT DETECTED	
ADOENIT	NOT DETECTED	
ABSENT	NOT DETECTED	
ARSENT	NOT DETECTED	
ABOLIVI	NOT BETEOTED	
ABSENT	NEGATIVE	
ABSENT	NEGATIVE	
ABSENT	NEGATIVE	
ADOENT	NECATIVE	
ABSENT	NEGATIVE	
2.2	0.5	/hnf
2-3	U-0	/hpf
0-1	0-5	/hpf
• 1		, . · · · · · · · ·
ABSENT	0-2	/hpf
		•
ABSENT	NOT DETECTED	
ABSENT	NOT DETECTED	
EE\\\/	NOT DETECTED	
Γ⊏VV	NOI DETECTED	
ABSENT	NOT DETECTED	
ABSENT		
	SLIGHTLY HAZY 5.0 1.010 ABSENT ABSENT	SLIGHTLY HAZY 5.0 4.6 - 8.0 1.010 1.005 - 1.030 ABSENT NOT DETECTED ABSENT NOT DETECTED ABSENT NOT DETECTED ABSENT NEGATIVE ABSENT NEGATIVE ABSENT NEGATIVE ABSENT NEGATIVE ABSENT NEGATIVE ABSENT NEGATIVE ABSENT O-5 0-1 0-5 ABSENT 0-2 ABSENT NOT DETECTED ABSENT NOT DETECTED ABSENT NOT DETECTED FEW NOT DETECTED ABSENT NOT DETECTED ABSENT NOT DETECTED

Note:

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

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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 ***

Prabha

Dr. PRABHA ANAND MBBS, MD(Microbiology) CONSULTANT MICROBIOLOGISTS Reg No. WBMC 92308

Lab No. : SG2/23-03-2024/SR8904397



Lab No. : SG2/23-03-2024/SR8904397 Lab Add.

Patient Name : CHUNKU BHUTIA Ref Dr. : Dr.MEDICAL OFFICER

Age : 34 Y 6 M 12 D Collection Date :

Gender : F Report Date : 23/Mar/2024 11:23AM



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 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 (79 mm). 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 112 mm. & Lt. kidney 106 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 (82 mm. x 42 mm). Endometrium (collapsed wall) is in midline. **Mid to lower body shows a subserous to intramural myoma measuring 41 x 30 mm at left lateral wall.** Cervix looks normal. Pouch of Douglas is free.

OVARIES

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

Right ovary measures 38 x 17 mm.

Left Ovary measures 33 x 23 mm.

IMPRESSION:

Uterine myoma.

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Patient Name : CHUNKU BHUTIA Ref Dr. : Dr.MEDICAL OFFICER

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Gender : F Report Date : 23/Mar/2024 11:23AM

DEPARTMENT OF ULTRASONOGRAPHY

Please correlate clinically.

Kindly note

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

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

Patient Identity not verified.

DR. Ziaul Mustafa

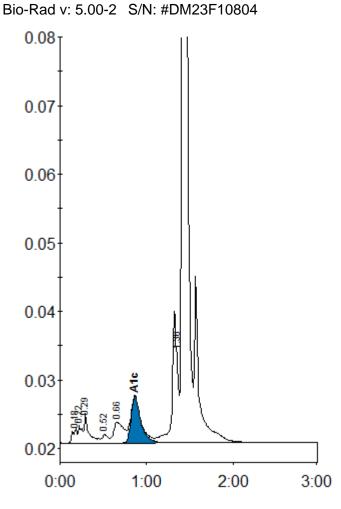
MD, Radiodiagnosis

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Patient report

Sample ID: D02132550156

Injection date 22/03/2024 05:23 PM
Injection #: 10 D-10 Method: HbA1c
Rack #: --- Rack position: 6



Peak table - ID: D02132550156

Peak	R.time	Height	Area	Area %
Unknown	0.18	2132	7510	0.5
A1a	0.22	2579	8196	0.5
A1b	0.29	4364	16583	1.1
F	0.52	1253	5985	0.4
LA1c/CHb-1	0.66	3031	24455	1.6
A1c	0.86	6809	53313	5.3
P3	1.36	19129	81713	5.4
A0	1.42	591938	1323606	87.0

Total Area: 1521359

Concentration:	%	mmol/mol
A1c	5.3	34