



Lab No. : BHT/13-01-2024/SR8628142	Lab Add. : Newtown,Kolkata-700156
Patient Name : GHULAM SIMNANI	Ref Dr. : Dr.MEDICAL OFFICER
Age : 36 Y 10 M 22 D	Collection Date : 13/Jan/2024 09:49AM
Gender : M	Report Date : 13/Jan/2024 06:44PM



DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
ALKALINE PHOSPHATASE , GEL SERUM (Method:IFCC standardization)	68	46-116	U/L
BILIRUBIN (DIRECT) (Method:Vanadate oxidation)	0.10	<0.2	mg/dL
SGOT/AST (Method:Modified IFCC)	24	13-40	U/L
SODIUM,BLOOD (Method:ISE INDIRECT)	141	132 - 146	mEq/L
CHLORIDE,BLOOD (Method:ISE INDIRECT)	104	99-109	mEq/L
CREATININE, BLOOD (Method:Jaffe, alkaline picrate, kinetic)	0.82	0.7-1.3	mg/dL
GLUCOSE,PP (Method:Gluc Oxidase Trinder)	98	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.

THYROID PANEL (T3, T4, TSH) , GEL SERUM			
T3-TOTAL (TRI IODOTHYRONINE) (Method:CLIA)	0.99	0.60-1.81 ng/ml	ng/ml
T4-TOTAL (THYROXINE) (Method:CLIA)	6.9	3.2-12.6	µg/dL
TSH (THYROID STIMULATING HORMONE) (Method:CLIA)	1.975	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:



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DEPARTMENT OF BIOCHEMISTRY

Test Name	Result	Bio Ref. Interval	Unit
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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)	23.5	19-49	mg/dL
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URIC ACID,BLOOD (Method:Uricase/Peroxidase)	5.70	3.5-7.2	mg/dL
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SGPT/ALT (Method:Modified IFCC)	28	7-40	U/L
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POTASSIUM,BLOOD (Method:ISE INDIRECT)	4.30	3.5-5.5	mEq/L
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BILIRUBIN (TOTAL) , GEL SERUM BILIRUBIN (TOTAL) (Method:Vanadate oxidation)	0.50	0.3-1.2	mg/dL
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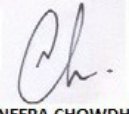
GLUCOSE,FASTING (Method:Gluc Oxidase Trinder)	83	Impaired Fasting-100-125 .-Diabetes- >= 126.-Fasting is defined as no caloric intake for at least 8 hours.	mg/dL
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In the absence of unequivocal hyperglycemia, diagnosis requires two abnormal test results 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.

*** End Of Report ***



Dr NEEPA CHOWDHURY
MBBS MD (Biochemistry)
Consultant Biochemist



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Patient Name : GHULAM SIMNANI	Ref Dr. : Dr.MEDICAL OFFICER
Age : 36 Y 10 M 22 D	Collection Date : 13/Jan/2024 09:49AM
Gender : M	Report Date : 13/Jan/2024 06:58PM

**DEPARTMENT OF BIOCHEMISTRY**

Test Name	Result	Bio Ref. Interval	Unit
PHOSPHORUS-INORGANIC,BLOOD (Method:Phosphomolybdate/UV) <i>ESTIMATED TWICE</i>	2.2	2.4-5.1 mg/dL	mg/dL

Suggested follow up

Correlate clinically

TOTAL PROTEIN [BLOOD] ALB:GLO RATIO , .			
TOTAL PROTEIN (Method:BIURET METHOD)	7.20	5.7-8.2 g/dL	g/dL
ALBUMIN (Method:BCG Dye Binding)	4.5	3.2-4.8 g/dL	g/dL
GLOBULIN (Method:Calculated)	2.70	1.8-3.2	g/dl
AG Ratio (Method:Calculated)	1.67	1.0 - 2.5	

LIPID PROFILE , GEL SERUM			
CHOLESTEROL-TOTAL (Method:Enzymatic)	188	Desirable: < 200 mg/dL Borderline high: 200-239 mg/dL High: > or =240 mg/dL	mg/dL
TRIGLYCERIDES (Method:GPO-Trinder)	123	Normal: < 150, BorderlineHigh::150-199, High:: 200-499, VeryHigh::>500	mg/dL
HDL CHOLESTEROL (Method:Elimination/catalase)	37	< 40 - Low 40-59- Optimum 60 - High	mg/dl
LDL CHOLESTEROL DIRECT (Method:Elimination / Catalase)	141	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)	10	< 40 mg/dl	mg/dl
CHOL HDL Ratio (Method:Calculated)	5.1	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.

CALCIUM,BLOOD (Method:Arsenazo III)	9.10	8.7-10.4 mg/dL	mg/dL
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URIC ACID, URINE, SPOT URINE			
URIC ACID, SPOT URINE	35.00	37-92 mg/dL	mg/dL

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DEPARTMENT OF BIOCHEMISTRY

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

GLYCATED HAEMOGLOBIN (HBA1C) , EDTA WHOLE BLOOD			
GLYCATED HEMOGLOBIN (HBA1C)	5.3	***FOR BIOLOGICAL REFERENCE INTERVAL DETAILS , PLEASE REFER TO THE BELOW MENTIONED REMARKS/NOTE WITH ADDITIONAL CLINICAL INFORMATION ***	%
HbA1c (IFCC) (Method:HPLC)	35.0		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 glycosylated hemoglobin measurement: the position of the IFCC Working Group. Clin Chem Lab Med. 2007;45(8):1077-1080.

[PDF Attached](#)

*** End Of Report ***

Dr. SANCHAYAN SINHA
 MBBS, MD, DNB (BIOCHEMISTRY)
 CONSULTANT BIOCHEMIST



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Gender : M	Report Date : 13/Jan/2024 07:01PM



DEPARTMENT OF HAEMATOLOGY

Test Name	Result	Bio Ref. Interval	Unit
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CBC WITH PLATELET (THROMBOCYTE) COUNT , EDTA WHOLE BLOOD

HEMOGLOBIN (Method:PHOTOMETRIC)	15.8	13 - 17	g/dL
WBC (Method:DC detection method)	6.9	4 - 10	*10 ³ /μL
RBC (Method:DC detection method)	5.23	4.5 - 5.5	*10 ⁶ /μL
PLATELET (THROMBOCYTE) COUNT (Method:DC detection method/Microscopy)	231	150 - 450*10 ³	*10 ³ /μL

DIFFERENTIAL COUNT

NEUTROPHILS (Method:Flowcytometry/Microscopy)	62	40 - 80 %	%
LYMPHOCYTES (Method:Flowcytometry/Microscopy)	25	20 - 40 %	%
MONOCYTES (Method:Flowcytometry/Microscopy)	11	2 - 10 %	%
EOSINOPHILS (Method:Flowcytometry/Microscopy)	02	1 - 6 %	%
BASOPHILS (Method:Flowcytometry/Microscopy)	00	0-0.9%	%

CBC SUBGROUP

HEMATOCRIT / PCV (Method:Calculated)	46.2	40 - 50 %	%
MCV (Method:Calculated)	88.4	83 - 101 fl	fl
MCH (Method:Calculated)	30.2	27 - 32 pg	pg
MCHC (Method:Calculated)	34.1	31.5-34.5 gm/dl	gm/dl
RDW - RED CELL DISTRIBUTION WIDTH (Method:Calculated)	13.7	11.6-14%	%
PDW-PLATELET DISTRIBUTION WIDTH (Method:Calculated)	14.7	8.3 - 25 fL	fL
MPV-MEAN PLATELET VOLUME (Method:Calculated)	8.6	7.5 - 11.5 fl	

ESR (ERYTHROCYTE SEDIMENTATION RATE) , EDTA WHOLE BLOOD

1stHour (Method:Westergren)	03	0.00 - 20.00 mm/hr	mm/hr
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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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DEPARTMENT OF HAEMATOLOGY

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

*** End Of Report ***

Kaushik Dey

MD (PATHOLOGY)
CONSULTANT PATHOLOGIST

Lab No. : BHT/13-01-2024/SR8628142

Lab Add. :

Patient Name : GHULAM SIMNANI

Ref Dr. : Dr.MEDICAL OFFICER

Age : 36 Y 10 M 22 D

Collection Date :

Gender : M

Report Date : 13/Jan/2024 11:47AM



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.

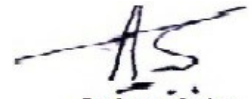
The cardio-thoracic ratio is normal.

Bony thorax reveals no definite abnormality.

IMPRESSION :

Normal study.

*** End Of Report ***


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



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DEPARTMENT OF CLINICAL PATHOLOGY

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))	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)	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	
<u>MICROSCOPIC EXAMINATION</u>			
LEUKOCYTES (PUS CELLS) (Method:Microscopy)	8-10	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

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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

MD (PATHOLOGY)
CONSULTANT PATHOLOGIST

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Ref Dr. : Dr.MEDICAL OFFICER

Age : 36 Y 10 M 22 D

Collection Date :

Gender : M

Report Date : 13/Jan/2024 11:18AM



DEPARTMENT OF CARDIOLOGY

REPORT OF E.C.G.

DATA		
HEART RATE	72	Bpm
PR INTERVAL	124	Ms
QRS DURATION	90	Ms
QT INTERVAL	354	Ms
QTC INTERVAL	393	Ms
AXIS		
P WAVE	59	Degree
QRS WAVE	69	Degree
T WAVE	51	Degree
IMPRESSION	:	Sinus rhythm. Normal ECG.

*** End Of Report ***

ACR

Dr. A C RAY
Department of Non-invasive
Cardiology

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Gender	: M	Report Date	: 13/Jan/2024 01:28PM



DEPARTMENT OF ULTRASONOGRAPHY
REPORT ON EXAMINATION OF WHOLE ABDOMEN

LIVER

Liver is normal in size having **early fatty changes seen**. 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 not dilated (0.47 cm.) with no intraluminal pathology (Calculi/mass) could be detected at its visualized part. Portal vein is normal (0.88 cm.) at porta.

GALL BLADDER

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

PANCREAS

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

SPLEEN

Spleen is normal in size (8.20 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 9.79 cm. & Lt. Kidney 8.85 cm.) axes & position. Cortical echogenicity appears normal maintaining cortico-medullary & cortico-hepatic differentiation. Margin is regular and cortical thickness is uniform. No calculus disease noted. No hydronephrosis changes detected. Visualized part of upper ureters are not dilated.

URINARY BLADDER

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

PROSTATE

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

It measures : 4.02 x 3.41 x 2.95 cm.

Approximate weight could be around = 21.18 gms.

RETROPERITONEUM & PERITONEUM

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

IMPRESSION

Early fatty changes in liver.

Kindly note

- Ultrasound is not the modality of choice to rule out subtle bowel lesion.

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- 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.

Nishan Ghosh

Dr. Nishan Ghosh
MBBS, CBET, FELLOW IN FOETAL
ECHOCARDIOGRAPHY FGI,
Reg.NO: 67862

Patient Data

Sample ID: D02135535366
 Patient ID: SR8628142
 Name:
 Physician:
 Sex:
 DOB:

Analysis Data

Analysis Performed: 01/13/2024 18:13:05
 Injection Number: 1202U
 Run Number: 33
 Rack ID: 0003
 Tube Number: 2
 Report Generated: 01/13/2024 18:27:04
 Operator ID: TRISHA

Comments:

Peak Name	NGSP %	Area %	Retention Time (min)	Peak Area
Unknown	---	0.1	0.110	3613
A1a	---	0.7	0.157	22223
A1b	---	1.0	0.221	29322
F	---	0.7	0.271	20065
LA1c	---	1.6	0.392	48956
A1c	5.3	---	0.495	134720
P3	---	3.2	0.786	98542
P4	---	1.2	0.861	36993
Ao	---	87.1	1.021	2667582

Total Area: 3,062,016

HbA1c (NGSP) = 5.3 % HbA1c (IFCC) = 35 mmol/mol

