



TEST REPORT

Reg. No :	2409100170
Name :	MR.ANAND BISOI
Age/Sex :	39 Years / Male
Ref. By :	MEDIWHEEL

UHID : UHID26713

 Reg. Date :
 07-Sep-2024

 Collected On :
 07-Sep-2024 09:46

 Report Date :
 07-Sep-2024

Parameter	<u>Result</u>	<u>Unit</u>	Reference Interval
COM	PLETE BL	OOD COUNT	Г (СВС)
Hemoglobin (SLS method)	14.4	g/dL	13.0 - 17.0
Hematrocrit (Electrical Impedance)	43.5	%	40 - 54
RBC Count (Electrical Impedance)	5.34	million/cmm	4.5 - 5.5
WBC Count (Flowcytometry)	9310	/cmm	4000 - 10000
Platelet Count (Electrical Impedance)	334000	/cmm	150000 - 410000
MCV (Calculated)	81.5	fL	83 - 101
MCH (Calculated)	27.0	Pg	27 - 32
MCHC (Calculated)	33.1	%	31.5 - 34.5
RDW (Calculated)	14.1	%	11.5 - 14.5
DIFFERENTIAL WBC COUNT			
Neutrophils (%)	58	%	38 - 70
Lymphocytes (%)	36	%	20 - 45
Monocytes (%)	05	%	2 - 8
Eosinophils (%)	01	%	1 - 4
Basophils (%)	00	%	0 - 1
Neutrophils (Absolute)	5400	/cmm	1800 - 7700
Lymphocytes (Absolute)	3300	/cmm	1000 - 3900
Monocytes (Absolute)	470	/cmm	200 - 800
Eosinophils (Absolute)	120	/cmm	20 - 500
Basophils (Absolute)	20	/cmm	0 - 100
Neutrophil-Lymphocyte Ratio(NLR)	1.63	/cmm	0.7 - 4.0
PERIPHERAL SMEAR EXAMINATION			
RBC Morphology	RBCs are No	ormochromic Norm	nocytic.
WBC Morphology	Total WBC a	nd differential cour	nt is within normal.
Platelets	Platelets are	adequate with nor	mal morphology.
Parasites	Malarial para	site is not detected	d.
ERYTHROCYTE SEDIMENTATION RAT	_		
ESR (After 1 hour)	12	mm/hr	0 - 14
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FBS Fasting Blood	č (<i>)</i>	101.3	mg/dL	70 - 110
Glucose Oxidase PPBS Post Prandial F	e-Peroxidase Blood Sugar (PPB	S) 129.0	mg/dL	110 - 140
Glucose Oxidas	0 (iiig/dE	

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Parameter		<u>Result</u>	Uni	<u>t Bic</u>	ological Reference Interval
	H		OBIN A1 C E		<u>N</u>
	H		OBIN A1 C E Specimen: Blood El		<u>N</u>
Hb A1C HPLC, NGSP Certifie				DTA >8 7-8 <7 6-7	Action Suggested , : Good Control , : Goal , : Near Normal Glycemia, : Non-diabetic Level

1. HbA1c >/= 6.5 *Or

2. Fasting plasma glucose >126 gm/dL. Fasting is defined as no caloric intake at least for 8 hrs.Or

3. Two hour plasma glucose >/= 200mg/dL during an oral glucose tolerence test by using a glucose load containing equivalent of 75 gm anhydrous glucosedissolved in water.Or

4. In a patient with classic symptoms of hyperglycemia or hyperglycemic crisis, a random plasma glucose >/= 200 mg/dL. *In the absence of unequivocal hyperglycemia, criteria 1-3 should be confirmed by repeat testing. American diabetes association. Standards of medical care in diabetes 2011. Diabetes care 2011;34;S11.

Importance of HbA1C (Glycated Hb.) in Diabetes Mellitus:

- HbA1C, also known as glycated heamoglobin, is the most important test for the assessment of long term blood glucose control(also called glycemic control).

- HbA1C reflects mean glucose concentration over pas 6-8 weeks and provides a much better indication of longterm glycemic control than blood glucose determination.

- HbA1c is formed by non-enzymatic reaction between glucose and Hb. This reaction is irreversible and therefore remains unaffected by short term fluctuations in blood glucose levels.

- Long term complications of diabetes such as retinopathy (Eye-complications), nephropathy (kidney-complications) and neuropathy (nerve complications), are potentially serious and can lead to blindness, kidney failure, etc.- Glyemic control monitored by HbA1c measurement using HPLC method (GOLD STANDARD) is considered most important. (Ref. National Glycohaemoglobin Standardization Program - NGSP).

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Parameter	<u>Result</u>	<u>Unit</u>	Reference Interval
	LIVER FUN	ICTION TES	<u>T</u>
SGPT	64.0	U/L	1 - 45
Optimized UV-IFCC			
SGOT	24.5	U/L	1 - 35
Optimized UV-IFCC			
Total Bilirubin	0.42	mg/dL	0 - 2.0
DCA method			
Direct Bilirubin	0.15	mg/dL	0.0 - 0.4
DCA method			
INDIRECT BILIRUBIN	0.27	mg/dL	0.0 - 1.6
Calculated			
Alkaline Phosphatase	54.3	U/L	53 - 128
PNP-AMP Buffer, Multiple-point rate			
Total Protein	6.43	g/dL	6.4 - 8.2
Albumin	4.12	g/dL	3.5 - 5.2
By Bromocresol Green			
Globulin	2.31	g/dL	2.3 - 3.5
Calculated			
A/G Ratio	1.78		0.8 - 2.0
Calculated			
GGT	27.3	U/L	1 - 55
HBsAg	Non - Reactive		
Immunochromatography			

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Parameter	<u>Result</u>	<u>Unit</u>	Reference Interval	
	RENAL F	UNCTION T	<u>EST</u>	
Creatinine	0.85	mg/dL	0.7 - 1.3	
Enzymatic ,IDMS Traceable				
Urea	25.3	mg/dL	19.0 - 45.0	
Urease-GLDH, enzmatic UV				
BUN	11.82	mg/dL	7 - 18	
Calculated				
Uric Acid	4.2	mg/dL	3.5 - 7.2	
Enzymatic using TBHBA				
Sodium	144.1	mmol/L	137 - 145	
Direct ISE				
Potassium	4.78	mmol/L	3.6 - 5.1	
Direct ISE				
Chloride	95.3	mmol/L	94 - 110	
Direct ISE				
Ionized Calcium	4.98	mg/dL	4.4 - 5.4	
Direct ISE				

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Parameter	<u>Result</u>	<u>Unit</u>	Biological Reference Interval
	LIP	ID PROFILE	
Cholesterol CHOD-PAP method	169	mg/dL	Desirable : < 200.0 Borderline High : 200-239 High : > 240.0
Triglyceride Enzymatic with GPO method	235.3	mg/dL	Normal : < 150.0 Borderline : 150-199 High : 200-499 Very High : > 500.0
VLDL Calculated	47.06	mg/dL	15 - 35
LDL CHOLESTEROL	88.04	mg/dL	Optimal : < 100.0 Near / above optimal : 100-129 Borderline High : 130-159 High : 160-189 Very High : >190.0
HDL Cholesterol Magnetic Cholesterol Oxidase	33.9	mg/dL	Low : < 40 High : > 60
Cholesterol /HDL Ratio	4.99		0 - 5.0
LDL / HDL RATIO	2.60		0 - 3.5
Total Lipids ^{Calculated}	768.60		400 - 1000

Pre-analytical requirements for given tests are -Fasting status anywhere between 10-12 hours before collection. Avoid alcohol beverages before lipid panel - minimum 24 hrs.

Lipid profile results can be erroneous if pre-analytical requirements are not met properly.

Any medical decision based on test results is to be taken with 2 or more consecutive results suggesting • pattern.

Please note that any lipid lowering drug may interfere in results estimation.

Sudden commencement or sudden withdrawal of Lipid lowering drug will interfere with test result.

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WHERE CARE MEETS COMFORT

Parameter



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Result

	THYROII	D FUNCTION T	EST
T3 (Triiodothyronine) CMIA	1.12	ng/mL	0.6 - 1.81
T4 (Thyroxine)	4.89	µg/dL	4.5 - 12.5
TSH	3.120	µIU/mI	0.35 - 4.94

Unit

ELFA-Enzyme Linked Fluorescent Assay

Thyroid stimulating hormone (TSH) is synthesized and secreted by the anterior pituitary in response to a negative feedback mechanism involving concentrations of FT3 (free T3) and FT4 (free T4). Additionally, the hypothalamic tripeptide, thyrotropin-relasing hormone (TRH), directly stimulates TSH production. TSH stimulates thyroid cell production and hypertrophy, also stimulate the thyroid gland to synthesize and secrete T3 and T4.Quantification of TSH is significant to differentiate primary (thyroid) from secondary (pituitary) and tertiary(hypothalamus) hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiaryhypothyroidism, TSH levels are low.

TSH levels During Pregnancy :

First Trimester :0.1 to 2.5 µIÚ/mL

Second Trimester : 0.2 to 3.0 µIU/mL

Third trimester : 0.3 to 3.0 µIU/mL

Referance : Carl A.Burtis, Edward R.Ashwood, David E.Bruns. Tietz Textbook of Clinical Chemistry and Molecular Diagnostics. 5th

Eddition.

Philadelphia: WB Sounders, 2012:2170

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URINE ROUTINE EXAMINATION

PHYSICAL EXAMINATION		
Quantity	10 cc	
Colour	Pale Yellow	
Clarity	Clear	
CHEMICAL EXAMINATION (BY REFLECTANCE PHOTOMETRIC METHOD)		
рН	7.0	4.6 - 8.0
Sp. Gravity	1.015	1.002 - 1.03
Protein	Nil	
Glucose	Nil	
Ketone Bodies	Nil	
Urobilinogen	Nil	
Bilirubin	Nil	
Nitrite	Nil	
Leucocytes	Nil	
Blood	Nil	
MICROSCOPIC EXAMINATION (MANUAL BY MICROSCOPY)		
Leucocytes (Pus Cells)	1 - 5/hpf	
Erythrocytes (Red Cells)	Nil	
Epithelial Cells	1-2/hpf	
Amorphous Material	Nil	
Casts	Nil	
Crystals	Nil	
Bacteria	Nil	
Yeast	Nil	
T. Vaginalis	Nil	
Spermatozoa	Nil	
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