

EXAMINATION OF EYES :-(BY OPHTALMOLOGIST)

Patient Name .	Mr. Malu	endra laur	nan Da	ite 7/10/23
Sex/Age	/ N	1R No	••••	Employee Id
EXTERNAL EXAMIN	ATION			
SQUINT		No		
NYSTAGMUS		- 140		
COLOUR VISION			mal	
FUNDUS:(RE):-		Weel (LE):-	cellul	
INDIVIDUAL COLOU	R IDENTIFICATION			
DISTANT VISION:(RE	6 6	(LE):-	646	
NEAR VISION:(RE):-	TPG (SY/6 (LE):-	Hts ((PG)
NIGHT BLINDNESS				
DIGUE	SPH	CYL	AXIS	ADD
RIGHT	-			+1.50
LEFT		-		+1.50
REMARKS :-		dus -		MBBS,MS(Ophthalm plogist Reg. No. CGMC 621/2006
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Dr. Sweety Lath

BDS (Cosmetic Dental Surgeon)



Dr. Vivek Lath

Chief Dental Consultant BDS, MDS, Diplomate (WCOI, Japan) Professor, MCDRC - Durg Reg. No. CGDC/14/PG/45

• Consult for : Digital Dentistry • Fixed Teeth • RCT • Dental Implants • Gums Diseases • Dentures • Cosmetic Filling • Tooth Jewellery Digital OPG
 Braces Treatment
 Tooth Removal
 Kids Dental Treatment
 All Kind of Dental Surgeries

Mr. Mahendra Dewangan

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Mr. Mahandra Devangan

BP-160/100, P-78/100 H-155 c.m W+-SCKg

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: Mr. MAHENDRA KUMAR DEWANG

Age/Gender

: 46 Y Male

UHID/ MR No

: 7120

OP Visit No

: OPD-UNIT-II-2

Visit Date

: 07/10/2023

Reported On

: 12/10/2023 05:55PM

Sample Collected On: 07/10/2023 05:33PM Ref. Doctor

: SELF

Sponsor Name

HAEMATOLOGY

	IIALIIIAIOLOG	•	
Investigation	Observed Value	Unit B	iological Reference Interval
CBC - COMPLETE BLOOD COUNT			
Haemoglobin(HB) Method: CELL COUNTER	13.3	gm/dl	12 - 17
Erythrocyte (RBC) Count Method: CELL COUNTER	4.91	mill/cu.mm.	4.20 - 6.00
PCV (Packed Cell Volume) Method: CELL COUNTER	39.90	%	39 - 52
MCV (Mean Corpuscular Volume) Method: CELL COUNTER	81.3	fL	76.00 - 100
MCH (Mean Corpuscular Haemoglobin) Method: CELL COUNTER	27.1	pg	26 - 34
MCHC (Mean Corpuscular Hb Concn.) Method: CELL COUNTER	33.3	g/dl	32 - 35
RDW (Red Cell Distribution Width) Method: CELL COUNTER	17.3	%	11- 16
Total Leucocytes (WBC) Count Method: CELL COUNTER	6.08	cells/cumm	3.50 - 10.00
Neutrophils Method: CELL COUNTER	49	%	40.0 - 73.0
Lymphocytes Method: CELL COUNTER	38	%	15.0 - 45.0
Monocytes	05	%	4.0 - 12.0
Eosinophils Method: CELL COUNTER	08	%	1-6%
Basophils Method: CELL COUNTER	00	%	0.0 - 2.0

End of Report

Results are to be corelated clinically

Lab Technician / Technologist path

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HAEMATOLOGY

Investigation

Observed Value

Unit

Biological Reference Interval

Platelet Count

189

150-400 lacs/cu.mm

Method: CELL COUNTER

1. As per the recommendation of International council for Standardization in Hematology, the differential leucocyte counts are additionally being reported as absolute numbers of each cell in per unit volume of blood.

2. Test conducted on EDTA whole blood.

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BIO CHEMISTRY

DIO OTILIMIOTIE		
Observed Value	Unit	Biological Reference Interval
278.0	mg/dl	70-140
173.0	mg/dl	70 - 120
10	mg/dl	7 - 20
0.94	mg/dl	0.6-1.4
4.20	mg/dL	2.6 - 7.2
	Observed Value 278.0 173.0 10 0.94	 278.0 mg/dl 173.0 mg/dl 10 mg/dl 0.94 mg/dl

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BIO CHEMISTRY

	BIO CHEINIS I	C1	
Investigation	Observed Value	Unit	Biological Reference Interval
LIPID PROFILE TEST (PACKA	GE)		
Cholesterol - Total	210.0	mg/dl	Desirable: < 200 Borderline High: 200-239 High: >= 240
Triglycerides level	369.0	mg/dl	Normal: < 150 Borderline High: 150-199 Very High: >=500
Method: Spectrophotomatric			
HDL Cholesterol	39.0	mg/dl	Major risk factor for heart disease: < 40 Negative risk factor for heart disease :>60
Method: Spectrophotomatric			None None
LDL Cholesterol	97.20	mg/dl	Optimal:< 100 Near Optimal:100 – 129 Borderline High: 130-159 High: 160-189 Very High :>=190
Method: Spectrophotomatric VLDL Cholesterol	73.80	mg/dl	6 - 38
Total Cholesterol/HDL Ratio	5.38		3.5-5
Methode: Spectrophotometric			

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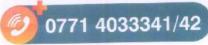
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BIO CHEMISTRY

*	BIO CHEIMISTR		
Investigation	Observed Value	Unit	Biological Reference Interval
LIVER FUNCTION TEST			
Bilirubin - Total	1.0	mg/dl	0.1- 1.2
Method: Spectrophotometric			
Bilirubin - Direct	0.2	mg/dl	0.05-0.3
Method: Spectrophotometric			
Bilirubin (Indirect)	0.80	mg/dl	0 - 1
Mathod: Calculated			
SGOT (AST)	40	U/L	0 - 40
Method: Spectrophotometric			
SGPT (ALT)	46	U/L	0 - 41
Method: Spectrophotometric			
ALKALINE PHOSPHATASE	63	U/L	25-147
Total Proteins	7.0	g/dl	6 - 8
Method: Spectrophotometric			
Albumin	4.8	mg/dl	3.4 - 5.0
Method: Spectrophotometric			
Globulin	2.2	g/dl	1.8 - 3.6
Mathod: Calculated			
A/G Ratio	2.18	%	1.1 - 2.2
Mathod: Calculated			

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BIO CHEMISTRY

Biological Reference Interval Unit Observed Value Investigation HbA1c (Glycosalated Haemoglobin) Non-diabeticc:<=5.6, Pre-

Diabetic 5.7-6.4, Diabetic:>=6.5

1.HbA1c is used for monitoring diabetic control. It reflects the estimated average glucose (eAG).

2.HbA1c has been endorsed by clinical groups & ADA (American Diabetes Association) guidelines 2017, for diagnosis of diabetes using a cut-off point of 6.5%.

Trends in HbA1c are a better indicator of diabetic control than a solitary test.

Low glycated haemoglobin(below 4%) in a non-diabetic individual are often associated with systemic inflam

1.HbA1c is used for monitoring diabetic control. It reflects the estimated average glucose (eAG).

2.HbA1c has been endorsed by clinical groups & ADA (American Diabetes Association) guidelines 2017, for diagnosis of diabetes using a cut-off point of 6.5%.

3. Trends in HbA1c are a better indicator of diabetic control than a solitary test.

4. Low glycated haemoglobin(below 4%) in a non-diabetic individual are often associated with systemic inflammatory diseases, chronic anaemia(especially severe iron deficiency & haemolytic), chronic renal failure and liver diseases. Clinical correlation suggested.

5. To estimate the eAG from the HbA1C value, the following equation is used: eAG(mg/dl) = 28.7*A1c-46.7

Interference of Haemoglobinopathies in HbA1c estimation.

A. For HbF > 25%, an alternate platform (Fructosamine) is recommended for testing of HbA1c.

B. Homozygous hemoglobinopathy is detected, fructosamine is recommended for monitoring diabetic status

C. Heterozygous state dete

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HAEMATOLOGY

Investigation

Observed Value

Unit

Biological Reference Interval

ESR- Erythrocyte Sedimentation Rate

15

mm /HR

0 - 10

Method: Westergren's Method

1. It indicates presence and intensity of an inflammatory process, never diagnostic of a specific disease. Changes are more significant than a single abnormal test.

2. It is a prognostic test and used to monitor the course or response to treatment of diseases like tuberculosis, bacterial endocarditis, acute rheumatic fever, rheumatoid arthritis, SLE, Hodgkins disease, temporal arteritis, polymyalgia rheumatica.

3. Also increased in pregnancy, multiple myeloma, menstruation & hypothyroidism

Blood Group (ABO Typing)

Blood Group (ABO Typing)

0

RhD factor (Rh Typing)

POSITIVE

End of Report

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IMMUNO ASSAY

Biological Reference Interval Unit Observed Value Investigation

PSA - TOTAL

PSA-TOTAL

1.210

ng/ml

Bordeerline: 4 - 10

10 - 49 years: 1.5 · 50 - 59 " : 2.5 60 - 69 " : 4.5

70 - 79": 7.5

- 1. PSA is detected in serum of males with normal, benign hypertrophic and malignant prostatitis.
- 2. Measurement of serum PSA level is not recommended as a screening procedure for the diagnosis of cancer, because elevated PSA levels also are observed in patients with benign prostatic hypertrophy.
- 3. The fact that PSA is unique to prostate tissue makes it a suitable marker for monitoring men with cancer of the prostate. PSA is also useful for determining possible recurrence after therapy when used in conjunction with other diagnostic indices.

METHOD: Fluorometric Immunoassay (Done with mini VIDAS Bio Meriux France)

PATHOLOGIST *All Reports Require Clinical Interpretation, please consult your Doctor

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IMMUNO ASSAY

Investigation Observed Value Unit Biological Reference Interval T3, T4, TSH

T3 (Total) by CLIA, serum

1.13

ng/mL

0.79 - 1.58

Clinical Use

· Diagnose and monitor treatment of Hyperthyroidism

Increased Levels: Pregnancy, Graves disease, T3 thyrotoxicosis, TSH dependent Hyperthyroidism,

Decreased Levels: Nonthyroidal illness, Hypothyroidism, Nutritional deficiency, Systemic illness, Decreased TBG

T4(Total) by CLIA, serum

6.30

mcg/dl

4.5-12.0

Clinical Use

· Diagnose Hypothyroidism and Hyperthyroidism when overt and / or due to pituitary or hypothalamic disease.

Increased Levels: Hyperthyroidism, Increased TBG, Familial dysalbuminemic hyperthyroxinemia, Increased Transthyretin, Estrogen therapy, Pregnancy

Decreased Levels: Primary hypothyroidism, Pituitary TSH deficiency, Hypothalamic TRH deficiency, Non thyroidal illness, Decreased TBG.

TSH (Ultrasensitive) CLIA Serum

3.75

mIU/ml

0.34 - 5.6

Initial test of thyroid function in patients with suspected thyroid dysfunction

· Assess thyroid status in patients with abnormal total T4 concentrations

· Distinguish Euthyroid hyperthyroxinemias from hypothyroidism. Increased Levels: Thyroid hormone resistance, Hyperthyroidism

Decreased Levels: Primary hypothyroidism, Secondary hypothyroidism

Clinical Use

· Initial test of thyroid function in patients with suspected thyroid dysfunction

Note: Total T3 & T4 levels measure the hormone which is in the bound form and is not available to most tissues. In addition severe systemic illness which affects the thyroid binding proteins can falsely alter Total T4 levels in the absence of a primary thyroid disease. Hence Free T3 & T4 levels are recommended for accurate assessment of thyroid dysfunction.

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CLINICAL PATHOLOGY

Investigation URINE ROUTINE EXAMINATION	Observed Value	Unit	Biological Reference Interval
Physical Examination			
Volum of urine	25ML		
Appearance	Clear		Clear
Colour	Pale Yellow		Colourless
Specific Gravity	1.010		1.001 - 1.030
Reaction (pH)	5.5		
Chemical Examination			
Protein(Albumin) Urine	Absent		Absent
Glucose(Sugar) Urine	Present 3 +		Absent
Blood	Absent		Absent
Leukocytes	Absent		Absent
Ketone Urine	Absent		Absent
Bilirubin Urine	Absent		Absent
Urobilinogen	Absent		Absent
Nitrite (Urine)	Absent		Absent
Microscopic Examination			
RBC (Urine)	NIL	/hpf	0 - 2
Pus cells	2-4	/hpf	0 - 5
Epithelial Cell	1 - 2	/hpf	0 - 5
Crystals	Not Seen	/hpf	Not Seen
Bacteria	Not Seen	/hpf	Not Seen
Budding yeast	Not Seen	/hpf	
	E 1 (B	3	

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