



# DIAGNOSTIC

RAIBARELI ROAD, TELIBAGH, LUCKNOW E-mail: mskdiagnosticspvt@gmail.com, Website: mskdiagnostics.in

Mobile: 7565000448

Collected At: (MSK)

Name : MR. JAGDAMBA PRASAD Ref/Reg No : 12913 / TPPC/MSK-Ref By

: Dr. MEDI WHEEL Sample : Blood, Urine

Age : 38 Yrs. Gender : Male

Registered Collected

: 14-1-2023 06:13 PM : 14-1-2023 08:36 AM

Received : 14-1-2023 06:13 PM Reported : 14-1-2023 06:16 PM

Investigation

Observed Values

Units

Biological Ref.

Interval

### **BIOCHEMISTRY**

### \*Glycosylated Hemoglobin (HbA1C)

\* Glycosylated Hemoglobin (HbA1C) 5.3 % 0-6 (Hplc method) Mean Blood Glucose (MBG) 111.38 mg/dl

: Non Diebetic Level

: Goal

> 8 % : Action suggested

SUMMARY

If HbAlc is >8% which causes high risk of developing long term complications like retinopathy, Nephropathy, Cardiopathy and Neuropathy. In diabetes mellitus sugar (glucose) accumulates in blood stream beyond normal level. Measurement of blood / plasma glucose level (in fasting, "after meal" i.e. PP or random condition) reflect acute changes related to immediate past condition of the patient which may be affected by factor like duration of fasting or time of intake of food before fasting, dosages of anti diabetic drugs, mental conditions like stress, anxiety etc. it does not indicate the long-term aspects of diabetic control.

Glucose combines with hemoglobin (Hb) continuously and nearly irreversibly during life span of RBC (120 days), thus glycosylated Hb is proportional to mean plasma glucose level during the previous 2-3 months. HBA1C, a glycosylated Hb comprising 3% - 6% of the total Hb in healthy may double of even triple in diabetes mellitus depending on the level of hyperglycemia(high blood glucose level), thus correlating with lack of control by monitoring diabetic patients compliance with therapeutic regimen used and long term blood glucose level control. Added advantage is its ability to predict progression of diabetic complications. HbAlc value is no way concerned with the blood sugar on the day of testing and dietary preparation of fasting is unnecessary.

DR. POONAM SINGH MD (PATH)

-- End of report --(SENIOR MECHNOLOGIST) (CHECKED BY)

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#### 2V

Plasma Glucose Fasting [Method: Hexokinase]	98.4	mg/dL	70 - 110
Plasma Glucose, PP (2 Hrs after meal) [Method: Hexokinase]	137.0	mg/dL.	120-170
Serum Bilirubin (Total)	0.4	mg/dl.	0.0 - 1.2
* Serum Bilirubin (Direct)	0.2	mg/dl.	0- 0.4
* Serum Bilirubin (Indirect)	0.2	mg/dl.	0.2-0.7
SGPT [Method: IFCC (UV without pyridoxal-5-phosphate]	18.2	IU/L	10 - 50
SGOT [Method: IFCC (UV without pyridoxal-5-phosphate]	20.9	IU/L	10 - 50
Serum Alkaline Phosphatase [Method:4-Nitrophenyl phosphate (pNPP)]	.139.0	IU/L	108 - 306
Serum Protein	7.7	gm/dL	6.2 - 7.8
Serum Albumin	4.8	gm/dL.	3.5 - 5.2
Serum Globulin	2.9	gm/dL.	2.5-5.0
A.G. Ratio	1.66:1		
* Gamma-Glutamyl Transferase (GGT)	25.4	IU/L	Less than 55

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### **HEMATOLOGY**

laemoglobin Method: SLSI	14.3	g/dL	13 - 17
HCT/PCV (Hematocrit/Packed Cell Volume)	43.8	ml %	36 - 46
Method: Derived] BC Count			
Method: Electrical Impedence]	5.21	10^6/µl	4.5 - 5.5
ICV (Mean Corpuscular Volume)	93.9	fL.	83 - 101
Method: Calculated]			
ICH (Mean Corpuscular Haemoglobin)	27.4	pg	27 - 32
Method: Calculated] MCHC (Mean Corpuscular Hb Concentration)	29.2	g/dL	31.5 - 34.5
Method: Calculated]		8/ 42	31.5 34.5
LC (Total Leucocyte Count)	12.2	10^3/μΙ	4.0 - 10.0
Method: Flow Cytometry/Microscopic] LC (Differential Leucocyte Count):			
Method: Flow Cytometry/Microscopic]			
olymorphs	78	%	40.0 - 80.0
ymphocytes	20	%	20.0 - 40.0
osinophils	02	%	1.0 - 6.0
Monocytes	00	%	2.0 - 10.0
latelet Count	290	10^3/μΙ	150 - 400
Method: Electrical impedence/Microscopic]			

*Observed Reading	18	mm for 1 hr	0-10
*Erythrocyte Sedimentation Rate (E.S.R.) [Method: Wintrobe Method]			

* ABO Typing	" B "	
* Rh (Anti - D)	Positive	

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Age

LIPID PROFILE (F)	The second secon	No. and	
Serum Cholesterol	148.0	mg/dL.	<200
Serum Triglycerides	77.5	mg/dL.	<150
HDL Cholesterol	35.6	mg/dL	>55
LDL Cholesterol	97	mg/dL.	<130
VLDL Cholesterol	16	mg/dL.	10 - 40 '
CHOL/HDL	4.16		
LDL/HDL	2.72	NOT ANY	

#### INTERPRETATION:

National Cholestrol Education program Expert Panel (NCEP) for Cholestrol:

Desirable : < 200 mg/dl
Borderline High : 200-239 mg/dl
High : =>240 mg/dl

National Cholestrol Education program Expert Panel (NCEP) for Triglycerides:

Desirable : < 150 mg/dl
Borderline High : 150-199 mg/dl
High : 200-499 mg/dl
Very High : >500 mg/dl

National Cholestrol Education program Expert Panel (NCEP) for HDL-Cholestrol:
<40 mg/dl : Low HDL-Cholestrol [Major risk factor for CHD]
=>60 mg/dl : Hight HDL-Cholestrol [Negative risk factor for CHD]

National Cholestrol Education program Expert Panel (NCEP) for LDL-Cholestrol:

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

[Method for Cholestrol Total: Enzymatic (CHOD/POD)]
[Method for Triglycerides: Enzymatic (Lipase/GK/GPO/POD)]

[Method for HDL Cholestrol: Homogenous Enzymatic (PEG Cholestrol esterase)]
[Method for LDL Cholestrol: Homogenous Enzymatic (PEG Cholestrol esterase)]

[Method for VLDL Cholestrol: Friedewald equation]

[Method for CHOL/HDL ratio: Calculated]
[Method for LDL/HDL ratio: Calculated]

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## HORMONE & IMMUNOLOGY ASSAY

Serum T3	2.01	ng/dl	0.846 - 2.02
Serum T4	14.63	ug/dl	5.13 - 14.06
Serum Thyroid Stimulating Harmone (T.S.H.) [Method: Flectro Chemiluminescence Immunoassay (F	1.40	uIU/mI	0.39 - 5.60

### SUMMARY OF THE TEST

- 1) Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values along with depressed TSH levels.
- 2) primary hypothyroidism is accompanied by depressed serum T3 and T4 values and elevated serum TSH levels.
- 3) Normal T4 levels accompanied by high T3 levels are seen in patients with T3
- thyrotoxicosis. 4) Slightly elevated T3 levels may be found in pregnancy and esterogen therapy, while depressed levels maybe encountered in severe illness, malnutrition, renalfailure and during therapy with drugs like propanlol and propylthiouracil.

  5) Elevated TSH levels may also be indicative of TSH secreting pituitary tumour.

Chart of normal thyroid TSH levels during first, second and third trimester of pregnancy .

Normal TSH Level Stage

0.1-2.5 ulU/ml First Trimester 0.2-3.0 ulU/ml Second Trimester 0.3 - 3.5 ulU/mlThird Trimester

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## **BIOCHEMISTRY**

KIDNEY FUNCTION TEST			
Blood Urea	20.9		
Serum Creatinine		mg/dL.	20-40
	0.51	mg/dL.	0.50 - 1.40
Serum Sodium (Na+)	138	mmol/L	135 - 150
Serum Potassium (K+)	5.9		
Serum Uric Acid		mmol/L	3.5 - 5.3
Serum one Acid	9.6	mg/dL.	3.4 - 7.0

[Method for Urea: UREASE with GLDH]

[Method for Creatinine: Jaffes/Enzymatic]

[Method for Sodium/Potassium: Ion selective electrode direct]

[Method for Uric Acid: Enzymatic-URICASE]

Serum Urea Blood Urea Nitrogen ( BUN ) 20.9

mg/dL.

10-45 6 - 21

### **CLINICAL PATHOLOGY**

Urine for Sugar (F)

Absent

Urine for Sugar (PP)

Absent

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PATIENT NAME:	MR. JAGDAMBA		
	PRASAD	AGE / SEX:	38Y/M
REF. BY :	MEDI WHEEL	DATE :	14.01.2023
			14.01.2023

# <u>USG – WHOLE ABDOMEN</u>

- Liver appears normal in shape, mildly enlarged in size (measuring ~17.87cm) & bright in echotexture with obscuring of vessels margins-suggestive of grade II fatty changes. No evidence of focal lesion is seen. No evidence of dilated IHBR seen. Portal vein appears normal in caliber.
- · CBD appears normal in caliber.
- Gall Bladder appears well distended with normal wall thickness. No calculus or changes of cholecystitis seen.
- Spleen is normal in shape, size (measuring ~12.69cm) and echotexture with no focal lesion within.
- Pancreas appears normal in size, shape &echopattern.
- Para-aortic region appears normal with no e/o lymphadenopathy.
- Right kidney measuring ~11.35cm; Left kidney measuring ~10.87cm. Both kidneys appear normal in position, shape, size & echotexture. CMD is normal. No calculus or hydronephrosis on either side.
- Urinary bladder appears well distended with no calculus or mass within.
- **Prostate** appears normal in size (~13cc) & echotexture.
- No evidence of ascites or pleural effusion seen.
- No abnormal bowel wall thickening or significant abdominal lymphadenopathy is seen.

## **IMPRESSION:**

Mild hepatomegaly with grade II fatty changes in liver. No focal parenchymal lesion.

Dr. Sarvesh Chandra Mishra

M.D., DNB Radio-diagnosis

PDCC Neuroradiology (SGPGI, LKO)

Ex- senior Resident (SGPGI, LKO)

European Diploma in radiology (EDiR), DICRI

Dr. Sweta Kumari

MBBS, DMRD

DNB Radio Diagnosis

Ex- Senior Resident Apollo Hospital Bengaluru

Ex- Resident JIPMER, Pondicherry

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### **CLINICAL PATHOLOGY**

#### URINE EXAMINATION ROUTINE

[Method: Visual, Urometer-120, Microscopy]

Physical Examination

Color

Volume

Sample

Investigation

Light Yellow

30

mL

RBC/µl

**Chemical Findings** 

Blood Bilirubin Urobilinogen

Ketones **Proteins Nitrites** Glucose

pH Specific Gravity

Leucocytes

Absent

Absent Absent

Absent Absent Absent

Absent 5.5 1.025

Absent

WBC/µL

Absent Absent

Absent

Absent Absent Absent

Absent 5.0 - 9.0

1.010 - 1.030 Absent

### Microscopic Findings

Red Blood cells Pus cells **Epithelial Cells** Casts Crystals

Amorphous deposit Yeast cells

Bacteria Others

Absent Occasional

Absent **Absent** Absent

Absent Absent

Absent

Absent

/HPF /HPF /HPF

/HPF /HPF /HPF /HPF

/HPF

/HPF

0-3 Absent/Few Absent Absent

Absent

Absent Absent

Absent Absent

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