

MSK

(A Complete Diagnostic Pathology Laboratory)

DIAGNOSTICS

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

Mobile: 7565000448

Collected At : JAVITRI

Name: MRS. NIRMALA KUMARI

Ref/Reg No : 107007 / TPPC\JAV-

Ref By : D Sample : B

: Dr. MEDI WHEEL : Blood, Urine

Sample(s)

: Plain, EDTA, Urine, FBS, PPP

Age : 44 Yrs. Gender : Female Registered

istered : 11-3-2023 03:24 PM

Collected

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: 11-3-2023 03:24 PM

Received Reported

: 12-3-2023 05:01 PM

Investigation	Observed Values	Units	Biological Ref Interval
HEMOGRAM			
(Method: Electrical impedance, Flowcytometry, Sepct	rophotometry)		
Haemoglobin	9.8	g/dL	11.5 - 15
[Method: SLS] HCT/PCV (Hematocrit/Packed Cell Volume)	20	10/	26.46
[Method: Derived]	30	ml %	36 - 46
RBC Count	3.77	10^6/μl	3.8 - 4.8
[Method: Electrical Impedence]		•	
MCV (Mean Corpuscular Volume) [Method: Calculated]	82.5	fL.	83 - 101
MCH (Mean Corpuscular Haemoglobin)	26.0	pg	27 - 32
[Method: Calculated]	30.0	Pb	27 32
MCHC (Mean Corpuscular Hb Concentration)	31.5	g/dL	31.5 - 34.5
[Method: Calculated]	6.6	4042/	
TLC (Total Leucocyte Count) [Method: Flow Cytometry/Microscopic]	6.6	10^3/µl	4.0 - 10.0
DLC (Differential Leucocyte Count):			
Method: Flow Cytometry/Microscopic]			
Polymorphs	68	%	40.0 - 80.0
Lymphocytes	29	%	20.0 - 40.0
Eosinophils	01	%	1.0 - 6.0
Monocytes	02	%	2.0 - 10.0
Platelet Count	191	10^3/μl	150 - 400
Method: Electrical impedence/Microscopic		== = / F **	223 100

*Erythrocyte Sedimentation Rate (E.S.R [Method: Wintrobe Method]	.)		
*Observed Reading	26	mm for 1 hr	0-20
* ABO Typing	"А"		
* Rh (Anti - D)	Positive		

Checked by

DR. MINAKSHI KAR ed." (MD PATH & BACT)

"The results generated here is subjected to the sample submitted."

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Investigation	Observed Values	Units	Biological Ref. Interval
Plasma Glucose Fasting	116	mg/dL	70 - 110
Plasma Glucose PP(2 Hrs after meal) [Method: Hexokinase]	126	mg/dL.	110-170
Glycosylated Hemoglobin (HbA1C) (Hplc method)	6.5	%	0 - 6
Mean Blood Glucose (MBG)	140	mg/dl	

Age

Gender : Female

SUMMARY

< 6 % : Non Diebetic Level

6-7 % : Goal

: Action suggested

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.

----- End of report -----

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[Method: Calculated]

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A.G. Ratio

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Investigation	Observed Values	Units	Biological Ref Interval
LIVER FUNCTION TEST			
Serum Bilirubin (Total)	0.27	mg/dl.	0.0 - 1.2
* Serum Bilirubin (Direct)	0.10	mg/dl.	0-0.4
* Serum Bilirubin (Indirect)	0.17	mg/dl.	0.2-0.7
Serum Alkaline Phosphatase	120	IU/L	35-104
Method:4-Nitrophenyl phosphate (pNPP)] GPT	16.0	IU/L	10-50
Method: IFCC (UV without pyridoxal-5-phosphate] GOT	18	IU/L	10-50
Method: IFCC (UV without pyridoxal-5-phosphate] Gamma-Glutamyl Transferase (GGT)	14.93	IU/L	Less than 38
erum Protein	6.9	gm/dL	6.2 - 7.8
Method: Biuret) erum Albumin Method: BCG)	4.5	gm/dL.	3.5 - 5.2
erum Globalin	2.4	gm/dL.	2.5-5.0

KIDNEY FUNCTION TEST			
Serum Urea	22.5	mg/dL.	10-45
Blood Urea Nitrogen (BUN)	11.02	mg/dL.	6 - 21
Serum Creatinine [Method: Jaffes Method/Enzymatic]	0.43	mg/dL.	0.40 - 1.00
Serum Sodium (Na+)	136	mmol/L	135 - 150
Serum Potassium (K+) [Method: Ion selective electrode direct]	3.90	mmol/L	3.5 - 5.5
Serum Uric Acid	4.40	mg/dL.	2.4 - 5.7
[Method for Uric Acid: Enzymatic-URICASE] * Serum Calcium (Total)	8.7	mg/dl.	8.2 - 10.2

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Ambulance Available

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Investigation	Observed Values	Units	Biological Ref. Interval
LIPID PROFILE			
Serum Cholesterol	193	mg/dL.	<200
Serum Triglycerides	87	mg/dL.	<150
HDL Cholesterol	47	mg/dL	>55
LDL Cholesterol	129	mg/dL.	<130
VLDL Cholesterol	17	mg/dL.	10 - 40
CHOL/HDL	4.11		
LDL/HDL	2.74		

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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Investigation Observed Values Units Biological Ref.

			intervai
T3, T4, TSH (ECLIA METHOD)			
	1 40	ng/dl	0.84 - 2.02
Serum T3	1.40	ng/dl	
Serum T4	9.35	ug/dl	5.13 - 14.6
Serum Thyroid Stimulating Harmone (T.S.H.) [Method: Electro Chemiluminescence Immunoassay (2.47 ECLIA)]	uIU/ml	0.39 - 5.60

SUMMARY OF THE TEST

- 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

Stage Normal TSH Level

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

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¹⁾ Primary hyperthyroidism is accompanied by elevated serum T3 and T4 values along with depressed TSH levels.



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Investigation

Observed Values

Gender : Female

Age

Units

mi

RBC/µl

WBC/µL

/HPF

/HPF

/HPF

Biological Ref.

Interval

URINE EXAMINATION ROUTINE

PHYSICAL EXAMINATION

Color Volume

CHEMICAL EXAMINATION Blood

Bilirubin Urobilinogen

Chyle [Method: Ether] Ketones

Nitrites Proteins

Glucose pН

Specific Gravity

Leucocytes MICROSCOPIC EXAMINATION

Red Blood cells

Pus cells **Epithelial Cells** Casts

Crystals Amorphous deposit Yeast cells

Bacteria **Parasites** Spermatozoa **Light Yellow**

25

Absent

Absent Absent

Absent

Absent Absent **Absent**

Absent 6.0 1.015

Absent

Absent Occasional 1-2

Absent Absent Absent Absent

Absent

Absent

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Absent

/HPF /HPF /HPF /HPF /HPF /HPF /HPF

Absent

Absent Absent Absent Absent

Absent

Absent

Absent

5.0 - 9.0

1.010 - 1.030 Absent

Absent

0-3 Absent/Few

Absent Absent Absent Absent

Absent Absent Absent

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Mobile: 7565000448

NAME: -MS. NIRMALA

DATE: -11.03.2023

REF.BY: - MEDIWHEEL

AGE: - 43Y/F

USG - ABDOMEN-PELVIS

Liver appears normal in shape, bulky in size (measuring ~14.96cm) & bright in echotexture without obscuration of vessels margins. 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 ~10.61cm) 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 ~12.03cm; Left kidney measuring ~11.60cm. 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.

Uterus is anteverted, normal in size, shape & echotexture.

Both ovaries appear normal. No evidence of adnexal mass on either side.

No free fluid in peritoneal cavity.

No abnormal bowel wall thickening or significant abdominal lymphadenopathy is seen.

IMPRESSION

• Bulky liver with grade I fatty changes. No focal parenchymal lesion is seen. Rest unremarkable USG abdomen-pelvis study.

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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DATE:-11/03/2023

REF.BY:-MEDI-WHEEL

AGE:-44Y/F

X-RAY CHEST (P.A. View)

- Lung fields are clear.
- No focal parenchymal lesion is noted.
- Mediastinum is central.
- Cardiac size is normal.
- C.P. angles are normally visualized.
- · Domes of diaphragm are normal.
- · Pulmonary hila appear normal.
- Soft tissue and bones are normal.

OPINION:

- No significant abnormality detected.
 - -Suggested clinical correlation.

Dr. Sarvesh Chandra Mishra

M.D., D.N.B. Radio-diagnosis PDCC Neuroradiology (SGPGIMS, LKO) Ex- senior Resident (SGPGIMS, LKO) European Diploma in radiology EDIR, DICRI Dr. Sweta Kumari

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