

Collected At : (MSK)

| | | |
|-------------------------------|-----------------|---------------------------------|
| Name : MRS. JYOTI MISHRA | Age : 38 Yrs. | Registered : 28-1-2023 11:28 AM |
| Ref/Reg No : 13058 /TPPC/MSK- | Gender : Female | Collected : 28-1-2023 09:10 AM |
| Ref By : Dr. MEDI WHEEL | | Received : 28-1-2023 11:28 AM |
| Sample : Blood, Urine | | Reported : 28-1-2023 05:19 PM |

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BIOCHEMISTRY

*Glycosylated Hemoglobin (HbA1C)

| | | | |
|--|--------|-------|-----|
| * Glycosylated Hemoglobin (HbA1C) (Hplc method) | 9.7 | % | 0-6 |
| * Mean Blood Glucose (MBG) | 268.02 | mg/dl | |

< 6 % : Non Diabetic Level
6-7 % : Goal
> 8 % : Action suggested

SUMMARY

If HbA1c 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 or 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. HbA1c 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)

(SENIOR TECHNOLOGIST)
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HEMATOLOGY

HEMOGRAM

| | | | |
|---|------|---------------------|-------------|
| Haemoglobin [Method: SLS] | 13.3 | g/dL | 11.5 - 15 |
| HCT/PCV (Hematocrit/Packed Cell Volume) [Method: Derived] | 42.0 | ml % | 36 - 46 |
| RBC Count [Method: Electrical Impedence] | 4.73 | 10 ⁶ /μl | 3.8 - 4.8 |
| MCV (Mean Corpuscular Volume) [Method: Calculated] | 92.3 | fL | 83 - 101 |
| MCH (Mean Corpuscular Haemoglobin) [Method: Calculated] | 28.2 | pg | 27 - 32 |
| MCHC (Mean Corpuscular Hb Concentration) [Method: Calculated] | 33.1 | g/dL | 31.5 - 34.5 |
| TLC (Total Leucocyte Count) [Method: Flow Cytometry/Microscopic] | 8.6 | 10 ³ /μl | 4.0 - 10.0 |
| DLC (Differential Leucocyte Count): [Method: Flow Cytometry/Microscopic] | | | |
| Polymorphs | 70 | % | 40.0 - 80.0 |
| Lymphocytes | 26 | % | 20.0 - 40.0 |
| Eosinophils | 02 | % | 1.0 - 6.0 |
| Monocytes | 02 | % | 2.0 - 10.0 |
| Platelet Count [Method: Electrical impedance/Microscopic] | 188 | 10 ³ /μl | 150 - 400 |

*Erythrocyte Sedimentation Rate (E.S.R.)

[Method: Wintrobe Method]

*Observed Reading

24

mm for 1 hr

0-20

* ABO Typing

" B "

* Rh (Anti - D)

Positive

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BIOCHEMISTRY

| | | | |
|---|----------|--------|--------------|
| Plasma Glucose Fasting [Method: Hexokinase] | 198.2 | mg/dL | 70 - 110 |
| Plasma Glucose, PP (2 Hrs after meal) [Method: Hexokinase] | 272.7 | mg/dL. | 120-170 |
| Serum Bilirubin (Total) | 0.5 | mg/dl. | 0.0 - 1.2 |
| * Serum Bilirubin (Direct) | 0.1 | mg/dl. | 0- 0.4 |
| * Serum Bilirubin (Indirect) | 0.4 | mg/dl. | 0.2-0.7 |
| SGPT [Method: IFCC (UV without pyridoxal-5-phosphate)] | 27.6 | IU/L | 10 - 50 |
| SGOT [Method: IFCC (UV without pyridoxal-5-phosphate)] | 26.9 | IU/L | 10 - 50 |
| Serum Alkaline Phosphatase [Method:4-Nitrophenyl phosphate (pNPP)] | 134.6 | IU/L | 108 - 306 |
| Serum Protein | 6.8 | gm/dL | 6.2 - 7.8 |
| Serum Albumin | 4.1 | gm/dL. | 3.5 - 5.2 |
| Serum Globulin | 2.7 | gm/dL. | 2.5-5.0 |
| A.G. Ratio | 1.52 : 1 | | |
| * Gamma-Glutamyl Transferase (GGT) | 32.2 | IU/L | Less than 38 |

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LIPID PROFILE (F)

| | | | |
|---------------------|-------|--------|---------|
| Serum Cholesterol | 169.9 | mg/dL. | <200 |
| Serum Triglycerides | 89.8 | mg/dL. | <150 |
| HDL Cholesterol | 50.8 | mg/dL. | >55 |
| LDL Cholesterol | 101 | mg/dL. | <130 |
| VLDL Cholesterol | 18 | mg/dL. | 10 - 40 |
| CHOL/HDL | 3.34 | | |
| LDL/HDL | 1.99 | | |

INTERPRETATION:

National Cholesterol Education program Expert Panel (NCEP) for Cholesterol:
Desirable : < 200 mg/dl
Borderline High : 200-239 mg/dl
High : =>240 mg/dl

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

National Cholesterol Education program Expert Panel (NCEP) for LDL-Cholesterol:
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 Cholesterol Total: Enzymatic (CHOD/POD)]
[Method for Triglycerides: Enzymatic (Lipase/GK/GPO/POD)]
[Method for HDL Cholesterol: Homogenous Enzymatic (PEG Cholesterol esterase)]
[Method for LDL Cholesterol: Homogenous Enzymatic (PEG Cholesterol esterase)]
[Method for VLDL Cholesterol: Friedewald equation]
[Method for CHOL/HDL ratio: Calculated]
[Method for LDL/HDL ratio: Calculated]

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BIOCHEMISTRY

| KIDNEY FUNCTION TEST | | | |
|----------------------|------|--------|-------------|
| Blood Urea | 19.7 | mg/dL. | 20-40 |
| Serum Creatinine | 0.60 | mg/dL. | 0.50 - 1.40 |
| Serum Sodium (Na+) | 142 | mmol/L | 135 - 150 |
| Serum Potassium (K+) | 4.8 | mmol/L | 3.5 - 5.3 |
| Serum Uric Acid | 3.0 | mg/dL. | 2.4 - 5.7 |

[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 | 19.7 | mg/dL. | 10-45 |
| Blood Urea Nitrogen (BUN) | 9.21 | mg/dL. | 6 - 21 |

CLINICAL PATHOLOGY

| | |
|---------------------|------------|
| Urine for Sugar (F) | Present ++ |
|---------------------|------------|

| | |
|----------------------|-------------|
| Urine for Sugar (PP) | Present +++ |
|----------------------|-------------|

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CLINICAL PATHOLOGY

URINE EXAMINATION ROUTINE

[Method: Visual, Urometer-120, Microscopy]

Physical Examination

| | | | |
|--------|-------------|----|--------------------|
| Color | Pale Yellow | | Light Yellow/Straw |
| Volume | 30 | mL | |

Chemical Findings

| | | | |
|------------------|------------|--------------|---------------|
| Blood | Absent | RBC/ μ l | Absent |
| Bilirubin | Absent | | Absent |
| Urobilinogen | Absent | | Absent |
| Ketones | Absent | | Absent |
| Proteins | Absent | | Absent |
| Nitrites | Absent | | Absent |
| Glucose | Present ++ | | Absent |
| pH | 6.0 | | 5.0 - 9.0 |
| Specific Gravity | 1.020 | | 1.010 - 1.030 |
| Leucocytes | Absent | WBC/ μ L | Absent |

Microscopic Findings

| | | | |
|-------------------|------------|------|------------|
| Red Blood cells | Absent | /HPF | Absent |
| Pus cells | Occasional | /HPF | 0-3 |
| Epithelial Cells | Absent | /HPF | Absent/Few |
| Casts | Absent | /HPF | Absent |
| Crystals | Absent | /HPF | Absent |
| Amorphous deposit | Absent | /HPF | Absent |
| Yeast cells | Absent | /HPF | Absent |
| Bacteria | Absent | /HPF | Absent |
| Others | Absent | /HPF | Absent |

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HORMONE & IMMUNOLOGY ASSAY

| | | | |
|---|------|--------|--------------|
| Serum T3 | 1.18 | ng/dl | 0.846 - 2.02 |
| Serum T4 | 6.59 | ug/dl | 5.13 - 14.06 |
| Serum Thyroid Stimulating Hormone (T.S.H.) | 1.78 | uIU/ml | 0.39 - 5.60 |
| [Method: Electro Chemiluminescence Immunoassay (ECLIA)] | | | |


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 estrogen therapy, while depressed levels maybe encountered in severe illness, malnutrition, renal failure and during therapy with drugs like propranolol and propylthiouracil.
- 5) Elevated TSH levels may also be indicative of TSH secreting pituitary tumour.

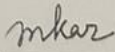
Chart of normal thyroid TSH levels during first, second and third trimester of pregnancy

| Stage | Normal TSH Level |
|------------------|------------------|
| First Trimester | 0.1-2.5 uIU/ml |
| Second Trimester | 0.2-3.0 uIU/ml |
| Third Trimester | 0.3-3.5 uIU/ml |

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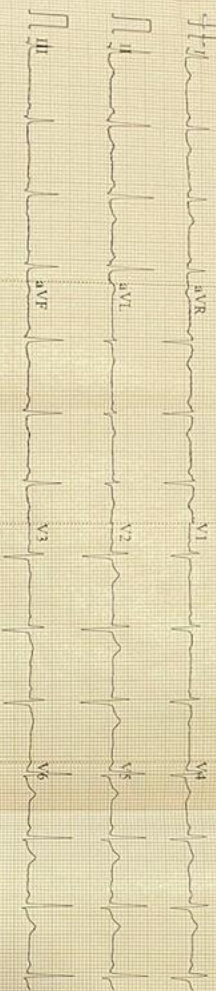
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ID: 496 28-01-2023 13:06:01



0.5-35Hz AC50 25mm/s 10mm/mV 81 V1.0 SEMIP V1.7 JAVTIRI HOSPITAL

ID: 496

Female
 Years () mmHg
 cm kg

Diagnosis-Information:

Sinus Rhythm
 Poor R Wave Progression(V3,V4)

HR 79 bpm
 P 160 ms
 PR 134 ms
 QRS 75 ms
 QT/QTc 348/401 ms
 P/QRST 42/63/27 °
 RV5/SV1 0.94/70.481 mV

Report Confirmed by:



MSK

(A Complete Diagnostic Pathology Laboratory)

DIAGNOSTICS

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

Mobile : 7565000448

NAME: - MS. JYOTI MISHRA

DATE: -28.01.2023

REF.BY: -MEDIWHEEL

AGE: - Y/F

USG – WHOLE ABDOMEN

Liver appears mildly enlarged in size (measures~ 169mm) shows diffusely increased echogenicity. No focal parenchymal lesion identified. No evidence of intra/ extrahepatic biliary tree dilatation noted. Portal vein appears to be of normal size.

Gall Bladder moderately distended. No definite calculi identified. No evidence of abnormal wall thickening noted.

Spleen appears normal in size (measures ~84mm), shape and echopattern No focal parenchymal identified.

Pancreas appears normal in size, shape and echopattern. No definite calcification or ductal dilatation noted.

Right kidney measures ~103x34mm; Left kidney measures ~106x52mm. Both kidneys appear normal in size, shape and echopattern. Corticomedullary differentiation appears maintained. No evidence of calculus or hydronephrosis on either side.

Urinary bladder appears partially distended. No obvious calculus or mass within.

Uterus anteverted appears normal in size measuring ~103x37mm. Myometrial echoes appears normal. The endometrial lining appears intact. Endometrial thickness measures ~ 8.5mm.

Right ovary measures~ 26x18mm; Left ovary measures ~ 28x13mm. Both ovaries appear normal in size, shape and echopattern

No evidence of ascites or pleural effusion seen. No significant retroperitoneal lymphadenopathy noted.

IMPRESSION:

- *Mild hepatomegaly with grade I fatty infiltration of liver.*
-Suggested clinical correlation

Dr. Sarvesh Chandra Mishra

M.D., DNB Radio-diagnosis
PDCC Neuroradiology (SGPGI, LKO)
Ex- senior Resident (SGPGI, LKO)
European Diploma in radiology EDiR, DICRI

Reports are subjected to human errors and not liable for medicolegal purpose.


Dr. Sweta Kumari

MBBS, DMRD
DNB Radio Diagnosis
Ex- Senior Resident Apollo Hospital Bengaluru
Ex- Resident JIPMER, Pondicherry

Reported by: Roli Vishvakarma

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