

Patient Name : MR. VYSHAK T V

Age / Gender : 36 years / Male

Patient ID : 53414

Referral : MediWheel

Collection Time : Aug 14, 2021, 08:49 a.m.

Reporting Time : Aug 14, 2021, 02:34 p.m.

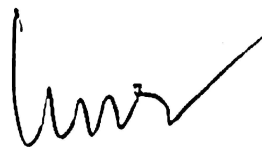
Sample ID :



003022621

Test Description	Value(s)	Reference Range	
<u>COMPLETE BLOOD COUNT (CBC)</u>			
Hemoglobin (Hb)	14.3	13.5 - 18.0	gm/dL
Erythrocyte (RBC) Count	4.86	4.7 - 6.0	mil/cu.mm
Packed Cell Volume (PCV)	41.4	42 - 52	%
Mean Cell Volume (MCV)	85.19	78 - 100	fL
Mean Cell Haemoglobin (MCH)	29.42	27 - 31	pg
Mean Corpuscular Hb Concn. (MCHC)	34.54	32 - 36	g/dL
Red Cell Distribution Width (RDW)	12.4	11.5 - 14.0	%
Total Leucocytes (WBC) Count	8600	4000-10000	cell/cu.mm
Neutrophils	56	40 - 80	%
Lymphocytes	34	20 - 40	%
Monocytes	5	2 - 10	%
Eosinophils	4	1 - 6	%
Basophils	1	1-2	%
Absolute Neutrophil Count	4816	2000 - 7000	/c.mm
Absolute Lymphocyte Count	2924	1000 - 3000	/c.mm
Absolute Monocyte Count	430	200 - 1000	/c.mm
Absolute Eosinophil Count	344	20 - 500	/c.mm
Absolute Basophils Count	86	20 - 100	/c.mm
Platelet Count	357	150 - 450	10 ³ /ul
Mean Platelet Volume (MPV)	9.0	7.2 - 11.7	fL
PCT	0.32	0.2 - 0.5	%
PDW	9.5	9.0 - 17.0	%

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URINE COMPLETE ANALYSIS,

Physical Examination

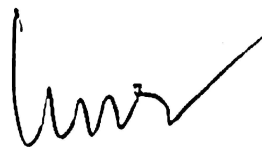
Quantity	26	-	ml
Colour	Pale Yellow	Pale yellow/Yellow	
Appearance	Clear	Clear	
Specific Gravity	1.015	1.005-1.025	
pH	6.0	5.0 - 8.0	
Deposit	Present	Absent	

Chemical Examination

Protein	Absent	Absent
Sugar	Absent	Absent
Ketones	Absent	Absent
Bile Salt	Absent	Absent
Bile Pigment	Absent	Absent
Urobilinogen	Normal	Normal

Microscopic Examination (/hpf)

Pus Cell	2-4	Upto 5
Epithelial Cells	1-2	Upto 5
Red Blood Cells	Absent	Absent
Casts	Absent	Absent
Crystals	Absent	Absent
Amorphous Deposit	Absent	Absent
Yeast Cells	Absent	Absent
Bacteria	Absent	Absent
Other findings	Not seen	Not seen



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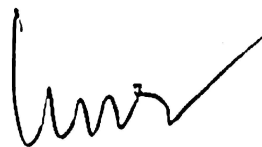
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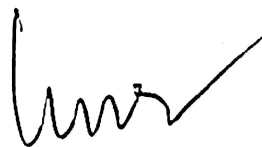
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Test Description	Value(s)	Reference Range
<u>BLOOD GROUP & RH TYPING</u>		
Blood Group (ABO typing) Method : Manual-Hemagglutination	"AB"	
RhD Factor (Rh Typing) Method : Manual hemagglutination	Positive	

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Test Description	Value(s)	Reference Range
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Glycosylated HbA1c

HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD	5.9	%
Method : (HPLC, NGSP certified)		
Estimated Average Glucose :	122.63	- mg/dL

Interpretation

As per American Diabetes Association (ADA)	
Reference Group	HbA1c in %
Non diabetic adults >=18 years	<5.7
At risk (Prediabetes)	5.7 - 6.4
Diagnosing Diabetes	>= 6.5
Therapeutic goals for glycemic control	Age > 19 years Goal of therapy: < 7.0 Action suggested: > 8.0 Age < 19 years Goal of therapy: <7.5

Note:

1. Since HbA1c reflects long term fluctuations in the blood glucose concentration, a diabetic patient who is recently under good control may still have a high concentration of HbA1c. Converse is true for a diabetic previously under good control but now poorly controlled .
2. Target goals of < 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targeting a goal of < 7.0 % may not be appropriate.

Comments

HbA1c provides an index of average blood glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control as compared to blood and urinary glucose determinations.



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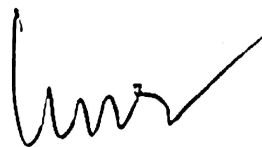
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ADA criteria for correlation between HbA1c & Mean plasma glucose levels.

HbA1c(%)	Mean Plasma Glucose (mg/dL)
6	126
7	154
8	183
9	212
10	240
11	269
12	298

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THYROID PROFILE TEST - TOTAL

T3-Total	85.63	60 - 200	ng/dL
T4-Total	7.30	4.52 - 12	ug/dL
TSH-Ultrasensitive	1.55	0.32 - 5.5	uIU/mL

Method : CLIA

Interpretation

TSH	T3	T4	Suggested Interpretation for the Thyroid Function Tests Pattern
Raised	Within range	Within range	Raised Within Range Within Range .Isolated High TSH especially in the range of 4.7 to 15 mIU/ml is commonly associated with Physiological & Biological TSH Variability. Subclinical Autoimmune Hypothyroidism. Intermittent 14 therapy for hypothyroidism .Recovery phase after Non-Thyroidal illness"
Raised	Decreased	Decreased	Chronic Autoimmune Thyroiditis Post thyroidectomy, Post radioiodine Hypothyroid phase of transient thyroiditis"
Raised or within range	Raised	Raised or within range	Interfering antibodies to thyroid hormones (anti-TPO antibodies) Intermittent 14 therapy or T4 overdose •Drug interference- Amiodarone, Heparin, Beta blockers, steroids, anti-epileptics.
Decreased	Raised or within range	Raised or within range	Isolated Low TSH -especially in the range of 0.1 to 0.4 often seen in elderly & Range Range associated with Non-Thyroidal illness .Subclinical Hyperthyroidism .Thyroxine ingestion'
Decreased	Decreased	Decreased	Central Hypothyroidism .Non-Thyroidal illness .Recent treatment for Hyperthyroidism (TSH remains suppressed)"
Decreased	Raised	Raised	Primary Hyperthyroidism (Graves' disease). Multinodular goitre, Toxic nodule •Transient thyroiditis: Postpartum, Silent (lymphocytic), Postviral (granulomatous, subacute, DeQuervain's), Gestational thyrotoxicosis with hyperemesis gravidarum"
Decreased Within Rang	Raised	Within range	T3 toxicosis •Non-Thyroidal illness
Within range	Decreased	Within range	Isolated Low T3-often seen in elderly & associated Non-Thyroidal illness In elderly the drop in T3 level can be upto 25%.

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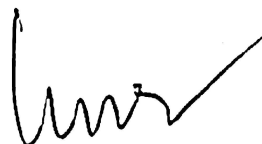
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Test Description	Value(s)	Reference Range	
<u>LIPID PROFILE</u>			
Cholesterol-Total Method : Spectrophotometry	223	Desirable level < 200 Borderline High 200-239 High >or = 240	mg/dL
Triglycerides Method : Serum, Enzymatic, endpoint	236	Normal: < 150 Borderline High: 150-199 High: 200-499 Very High: >= 500	mg/dL
HDL Cholesterol Method : Serum, Direct measure-PEG	36	Normal: > 40 Major Risk for Heart: < 40	mg/dL
LDL Cholesterol Method : Enzymatic selective protection	139.80	Optimal < 100 Near / Above Optimal 100-129 Borderline High 130-159 High 160-189 Very High >or = 190	mg/dL
VLDL Cholesterol Method : Serum, Enzymatic	47.20	6 - 38	mg/dL
CHOL/HDL Ratio Method : Serum, Enzymatic	6.19	3.5 - 5.0	
LDL/HDL Ratio Method : Serum, Enzymatic	3.88	2.5 - 3.5	

Note:

8-10 hours fasting sample is required.

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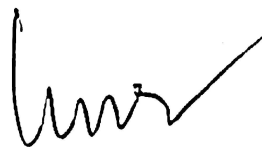
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Test Description	Value(s)	Reference Range	
<u>RENAL PROFILE</u>			
Urea Method : Uricase	23	19-42	mg/dL
Blood Urea Nitrogen-BUN Method : Serum, Urease	10.73	9-20	mg/dL
Creatinine Method : Serum, Jaffe	1.0	0.66-1.25	mg/dL
Uric Acid Method : Serum, Uricase	6.7	3.5-8.5	mg/dL
Sodium	152	137 - 145	mmol/L
Potassium	5.0	3.5 - 5.1	mmol/L
Chlorides	98	96.00 - 105.00	mmol/L

Remark:

In blood, Urea is usually reported as BUN and expressed in mg/dl. BUN mass units can be converted to urea mass units by multiplying by 2.14.

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Test Description	Value(s)	Reference Range	
<u>LIVER FUNCTION TEST</u>			
Total Protein Method : Serum, Biuret, reagent blank end point	8.8	6.3-8.2	g/dL
Albumin Method : Serum, Bromocresol green	5.1	3.5-5.0	g/dL
Globulin Method : Serum, EIA	3.70	1.8 - 3.6	g/dL
A/G Ratio Method : Serum, EIA	1.38	1.2 - 2.2	
Bilirubin - Total Method : Serum, Jendrassik Grof	0.7	0.3-1.2	mg/dL
Bilirubin - Direct Method : Serum, Diazotization	0.2	< 0.2	mg/dL
Bilirubin - Indirect Method : Serum, Calculated	0.50	0.1 - 1.0	mg/dL
SGOT Method : Serum, UV with P5P, IFCC 37 degree	77	17-59	U/L
SGPT Method : Serum, UV with P5P, IFCC 37 degree	140	21-72	U/L
Alkaline Phosphatase Method : PNPP-AMP Buffer/Kinetic	85	30 - 120	U/L
GGT-Gamma Glutamyl Transpeptidase Method : Serum, G-glutamyl-carboxy-nitroanilide	-	< 55	U/L

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Test Description	Value(s)	Reference Range
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PSA-Total (Prostate-specific antigen-Total)

PSA Profile *

PSA (Prostate Specific Antigen)-Total 0.75 0 - 4.0 ng/mL

Method : Serum, CLIA

Interpretation:

1. Increased levels are noted in Prostate cancer, Benign prostatic hypertrophy, Prostatitis

PSA (Prostate-Specific antigen)-Free * - 0.0 - 0.5 ng/mL

Method : Serum, CLIA

Interpretation & Remarks:


- Normal results do not eliminate the possibility of prostate cancer.
- Values obtained with different assay methods or kits may be different and cannot be used interchangeably.
- Tumor markers are not specific for malignancy. Test results cannot be interpreted as absolute evidence for the presence or absence of malignant disease
- Specimens drawn from patients undergoing prostate manipulation, especially needle biopsy and transurethral resection, may show erroneously high prostatic-specific antigen (PSA) results. Care should be taken that specimens are drawn before these procedures are performed.
- The percentage of free PSA can be used to estimate how likely it is that a biopsy will show cancer.
- If the percentage of free PSA is higher than 25%, the likelihood of prostate cancer is about 8%.
- If the percentage of free PSA is less than 10%, then the likelihood of prostate cancer rises to 56%.

Free PSA / Total PSA % - -

Method : Serum

Interpretation

- When total prostate-specific antigen (PSA) concentration is <2.0 ng/mL, the probability of prostate cancer in asymptomatic men is low, further testing and free PSA may provide little additional information. When total PSA concentration is >10.0 ng/mL, the probability of cancer is high and prostate biopsy is generally recommended.
- The total PSA range of 4.0 to 10.0 ng/mL has been described as a diagnostic "gray zone," in which the free:total PSA ratio helps to determine the relative risk of prostate cancer (see table below). Therefore, some urologists recommend using the free:total ratio to help select which men should undergo biopsy. However even a negative result of prostate biopsy does not



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Test Description	Value(s)	Reference Range
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rule-out prostate cancer. Up to 20% of men with negative biopsy results have subsequently been found to have cancer.

Based on free:total PSA ratio: the percent probability of finding prostate cancer on a needle biopsy by age in years:

Free:total PSA ratio	50-59 years	60-69 years	> or =70 years
< or =0.10	49.2%	57.5%	64.5%
0.11-0.18	26.9%	33.9%	40.8%
0.19-0.25	18.3%	23.9%	29.7%
>0.25	9.1%	12.2%	15.8%

Cautions

- Normal results do not eliminate the possibility of prostate cancer.
- Values obtained with different assay methods or kits may be different and cannot be used interchangeably
- Tumor markers are not specific for malignancy. Test results cannot be interpreted as absolute evidence for the presence or absence of malignant disease.

Interfering factors :

- Prostatic massage
- Proctoscopy
- Prostatic biopsy
- Prostate cancer patients receiving treatment with antiandrogens and luteinizing hormone-releasing factor agonists may exhibit markedly decreased levels of PSA. Also, men treated for benign prostatic hyperplasia with inhibitors of 5-alpha-reductase (finasteride) may demonstrate a significant reduction in PSA levels compared to values before treatment. Care should be taken in interpreting values for these individuals.
- In patients receiving therapy with high biotin doses (ie, >5 mg/day), no sample should be taken until at least 8 hours after the last biotin administration.

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Test Description	Value(s)	Reference Range
<u>GLUCOSE (F)</u>		
Glucose fasting Method : GOD-POD	100	Normal: 70 - 110 mg/dL

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Test Description	Value(s)	Reference Range
<u>GLUCOSE (PP)</u>		
Blood Glucose-Post Prandial Method : GOD-POD	127	70 - 120 mg/dL

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