

smart Health Report

An Insightful Health Analytics Report
for Easier Understanding



Prepared For

Mr DUMMY PL257M

M 25

Name
Mr DUMMY PL257M

Patient ID
10573467

Gender
M

Age
25

Health Summary



BLOOD COUNTS

Everything looks good



THYROID PROFILE

Everything looks good



LIPID PROFILE

Everything looks good



DIABETES MONITORING

Everything looks good



ANEMIA STUDIES

Everything looks good



VITAMIN PROFILE

Everything looks good



Patient NAME : Mr DUMMY PL257M
 DOB/Age/Gender : 25 Y/Male
 Patient ID / UHID : 10573467/OF10573467
 Referred BY : Self
 Sample Collected : Nov 26, 2024, 01:47 PM

Report STATUS : Final Report
 Barcode NO : HQ871336
 Sample Type : Whole blood EDTA
 Report Date : Nov 26, 2024, 02:45 PM.

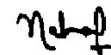


Test Description	Value(s)	Unit(s)	Reference Range
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Stress Impact- Male

Complete Blood Count (CBC)

RBC Parameters			
Hemoglobin <i>colorimetric</i>	15	g/dL	13.0 - 17.0
RBC Count <i>Electrical impedance</i>	4.9	10 ⁶ /μl	4.5 - 5.5
PCV <i>Calculated</i>	43.2	%	40 - 50
MCV <i>Calculated</i>	88.2	fl	83 - 101
MCH <i>Calculated</i>	30.6	pg	27 - 32
MCHC <i>Calculated</i>	34.5	g/dL	31.5 - 34.5
RDW (CV) <i>Calculated</i>	12.9	%	11.6 - 14.0
RDW-SD <i>Calculated</i>	37.3	fl	35.1 - 43.9
WBC Parameters			
TLC <i>Electrical impedance and microscopy</i>	5.1	10 ³ /μl	4 - 10
Differential Leucocyte Count			
Neutrophils <i>Laser based Flow-cytometry</i>	49	%	40-80
Lymphocytes <i>Laser based Flow-cytometry</i>	40	%	20-40
Monocytes <i>Laser based Flow-cytometry</i>	9	%	2-10
Eosinophils <i>Laser based Flow-cytometry</i>	2	%	1-6
Basophils <i>Laser based Flow-cytometry</i>	0	%	<2
Absolute Leukocyte Counts			
Neutrophils. <i>Calculated</i>	2.5	10 ³ /μl	2 - 7
Lymphocytes. <i>Calculated</i>	2.04	10 ³ /μl	1 - 3
Monocytes. <i>Calculated</i>	0.46	10 ³ /μl	0.2 - 1.0
Eosinophils. <i>Calculated</i>	0.1	10 ³ /μl	0.02 - 0.5
Basophils.	0	10 ³ /μl	0.02 - 0.5



Dr. Neha Prabhakar
MBBS, MD (Pathology)
Consultant Pathologist



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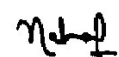
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Test Description	Value(s)	Unit(s)	Reference Range
<i>Calculated</i>			
Platelet Parameters			
Platelet Count <i>Electrical impedance and microscopy</i>	230	10 ³ /μl	150 - 410
Mean Platelet Volume (MPV) <i>Calculated</i>	9.2	fL	9.3 - 12.1
PCT <i>Calculated</i>	0.2	%	0.17 - 0.32
PDW <i>Calculated</i>	14.6	fL	8.3 - 25.0
P-LCR <i>Calculated</i>	26.8	%	18 - 50
P-LCC <i>Calculated</i>	62	10 ⁹ /L	44 - 140
Mentzer Index <i>Calculated</i>	18	%	> 13

Interpretation:
CBC provides information about red cells, white cells and platelets. Results are useful in the diagnosis of anemia, infections, leukemias, clotting disorders and many other medical conditions.



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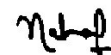
Erythrocyte Sedimentation Rate (ESR)

ESR - Erythrocyte Sedimentation Rate MODIFIED WESTERGREN	5	mm/hr	0 - 10
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Interpretation:

ESR is also known as Erythrocyte Sedimentation Rate. An ESR test is used to assess inflammation in the body. Many conditions can cause an abnormal ESR, so an ESR test is typically used with other tests to diagnose and monitor different diseases. An elevated ESR may occur in inflammatory conditions including infection, rheumatoid arthritis, systemic vasculitis, anemia, multiple myeloma, etc. Low levels are typically seen in congestive heart failure, polycythemia, sickle cell anemia, hypo fibrinogenemia, etc.

Reference- Dacie and lewis practical hematology



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Report STATUS : Final Report
 Barcode NO : HQ871336
 Sample Type : Whole blood EDTA
 Report Date : Nov 26, 2024, 04:41 PM.



Test Description	Value(s)	Unit(s)	Reference Range
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HbA1C (Glycosylated Haemoglobin)

Glycosylated Hemoglobin (HbA1c) <i>HPLC</i>	5.2	%	< 5.7
Estimated Average Glucose <i>Calculated</i>	102.54	mg/dL	Refer Table Below

Interpretation:

Interpretation For HbA1c% 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 Age < 19 years Goal of therapy: <7.5

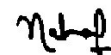
Note:

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

HbA1c(%)	Mean Plasma Glucose (mg/dL)	HbA1c(%)	Mean Plasma Glucose (mg/dL)
6	126	12	298
8	183	14	355
10	240	16	413



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Report STATUS : Final Report
 Barcode NO : ZF468879
 Sample Type : FLUORIDE F
 Report Date : Nov 27, 2024, 01:46 PM.



Test Description	Value(s)	Unit(s)	Reference Range
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Glucose Fasting (BSF)

Glucose Fasting <i>Hexokinase</i>	87.0	mg/dL	70 - 100
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Interpretation:

Status	Fasting plasma glucose in mg/dL
Normal	<100
Impaired fasting glucose	100 - 125
Diabetes	≥126

Reference : American Diabetes Association

Comment :

Blood glucose determinations are commonly used as an aid in the diagnosis and treatment of diabetes. Elevated glucose levels (hyperglycemia) may also occur with pancreatic neoplasm, hyperthyroidism, and adrenal cortical hyper function as well as other disorders. Decreased glucose levels (hypoglycemia) may result from excessive insulin therapy, insulinoma, or various liver diseases.

Note

- The diagnosis of Diabetes requires a fasting plasma glucose of $>$ or $=$ 126 mg/dL or a random / 2 hour plasma glucose value of $>$ or $=$ 200 mg/dL with symptoms of diabetes mellitus.
- Very high glucose levels ($>$ 450 mg/dL in adults) may result in Diabetic Ketoacidosis.



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 Sample Type : Serum
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Test Description	Value(s)	Unit(s)	Reference Range
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Lipid Profile

Total Cholesterol <i>Enzymatic - Cholesterol Oxidase</i>	126	mg/dL	<200
Triglycerides <i>Colorimetric - Lip/Glycerol Kinase</i>	89	mg/dL	<150
HDL Cholesterol <i>Phosphotungstic acid- Enzymatic</i>	47	mg/dL	> 40
Non HDL Cholesterol <i>Calculated</i>	79	mg/dL	<130
LDL Cholesterol <i>Calculated</i>	61.2	mg/dL	<100
V.L.D.L Cholesterol <i>Calculated</i>	17.8	mg/dL	< 30
Chol/HDL Ratio <i>Calculated</i>	2.68	Ratio	-
HDL/ LDL Ratio <i>Calculated</i>	0.77	Ratio	-
LDL/HDL Ratio <i>Calculated</i>	1.3	Ratio	-

Interpretation:

Lipid level assessments must be made following 9 to 12 hours of fasting, otherwise assay results might lead to erroneous interpretation. NCEP recommends of 3 different samples to be drawn at intervals of 1 week for harmonizing biological variables that might be encountered in single assays.

National Lipid Association Recommendations (NLA-2014)	Total Cholesterol (mg/dL)	Triglyceride (mg/dL)	LDL Cholesterol (mg/dL)	Non HDL Cholesterol (mg/dL)
Optimal	<200	<150	<100	<130
Above Optimal			100-129	130 - 159
Borderline High	200-239	150-199	130-159	160 - 189
High	>=240	200-499	160-189	190 - 219
Very High	-	>=500	>=190	>=220

HDL Cholesterol	
Low	High
<40	>=60

Risk Stratification for ASCVD (Atherosclerotic Cardiovascular Disease) by Lipid Association of India.

Risk Category	A. CAD with > 1 feature of high risk group
Extreme risk group	B. CAD with >1 feature of very high risk group of recurrent ACS (within 1 year) despite LDL-C <or = 50 mg/dl or poly vascular disease



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Test Description	Value(s)	Unit(s)	Reference Range
Very High Risk	1.Established ASCVD 2.Diabetes with 2 major risk factors of evidence of end organ damage 3. Familial Homozygous Hypercholesterolemia		
High Risk	1. Three major ASCVD risk factors 2. Diabetes with 1 major risk factor or no evidence of end organ damage 3. CHD stage 3B or 4. 4 LDL >190 mg/dl 5. Extreme of a single risk factor 6. Coronary Artery Calcium - CAC > 300 AU 7. Lipoprotein a >= 50 mg/dl 8. Non stenotic carotid plaque		
Moderate Risk	2 major ASCVD risk factors		
Low Risk	0-1 major ASCVD risk factors		
Major ASCVD (Atherosclerotic cardiovascular disease) Risk Factors			
1. Age >=45 years in Males & >= 55 years in Females	3. Current Cigarette smoking or tobacco use		
2. Family history of premature ASCVD	4. High blood pressure		
5. Low HDL			

Newer treatment goals and statin initiation thresholds based on the risk categories proposed by Lipid Association of India in 2020.

Risk Group	Treatment Goals		Consider Drug Therapy	
	LDL-C (mg/dl)	Non-HDL (mg/dl)	LDL-C (mg/dl)	Non-HDL (mg/dl)
Extreme Risk Group Category A	<50 (Optional goal <OR = 30)	<80 (Optional goal <OR = 60)	>OR = 50	>OR = 80
Extreme Risk Group Category B	>OR = 30	>OR = 60	> 30	> 60
Very High Risk	<50	<80	>OR = 50	>OR = 80
High Risk	<70	<100	>OR = 70	>OR = 100
Moderate Risk	<100	<130	>OR = 100	>OR = 130
Low Risk	<100	<130	>OR = 130*	>OR = 160

* After an adequate non-pharmacological intervention for at least 3 months.

References : Management of Dyslipidaemia for the Prevention of Stroke : Clinical practice Recommendations from the Lipid Association of India. Current Vascular Pharmacology,2022,20,134-155.



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High Sensitivity C-Reactive Protein (Hs-CRP)

HIGHLY SENSITIVE C-REACTIVE PROTEIN (hs-CRP) <i>immunoturbidimetric</i>	0.5	mg/L	< 1.00
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Interpretation:

Cardio CRP In mg/L	Cardiovascular Risk
<1	Low
1-3	Average
3-10	High
>10	Persistent elevation may represent Non cardiovascular inflammation

Note: To assess vascular risk, it is recommended to test hsCRP levels 2 or more weeks apart and calculate the average

Comments:

High sensitivity C Reactive Protein (hsCRP) significantly improves cardiovascular risk assessment as it is a strongest predictor of future coronary events. It reveals the risk of future Myocardial infarction and Stroke among healthy men and women, independent of traditional risk factors. It identifies patients at risk of first Myocardial infarction even with low to moderate lipid levels. The risk of recurrent cardiovascular events also correlates well with hsCRP levels. It is a powerful independent risk determinant in the prediction of incident Diabetes.



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Vitamin B12 / Cyanocobalamin

Vitamin - B12 ECLIA	357	pg/mL	-
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Interpretation:

Low Values are a sign of a vitamin B12 deficiency. People with this deficiency are likely to have or develop symptoms.

Causes of vitamin B12 deficiency include: Not enough vitamin B12 in diet (rare except with a strict vegetarian diet), Diseases that cause malabsorption (for example, celiac disease and Crohn's disease), Lack of intrinsic factor, Above normal heat production (for example, with hyperthyroidism), Pregnancy. Increased vitamin B12 levels are uncommon. Usually excess vitamin B12 is removed in the urine. Conditions that can increase B12 levels include: Liver disease (such as cirrhosis or hepatitis), Myeloproliferative disorders (for example, polycythemia vera and chronic myelocytic leukemia).

Vitamin B12: Low Levels can cause malabsorption, Lack of intrinsic factor, Above normal heat production (for example, with hyperthyroidism), Pregnancy. High Level Liver disease, Myeloproliferative disorders (for example, polycythemia vera and chronic myelocytic leukemia).

1. Out of 140 healthy indian population, 91% of Vitamin B 12 concentrations was at lower level: 59.00 pg/ml and upper level: 700.00 pg/ml

"Patients on Biotin supplement may have interference in some immunoassays. Ref: Arch Pathol Lab Med—Vol 141, November 2017. With individuals taking high dose Biotin (more than 5 mg per day) supplements, at least 8-hour wait time before blood draw is recommended."



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Vitamin D 25 Hydroxy

Vitamin D 25 - Hydroxy <i>ECLIA</i>	47	ng/mL	Deficiency : <30 ng/mL
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Interpretation:
 25-Hydroxy vitamin D represents the main body reservoir and transport form. Mild to moderate deficiency is associated with Osteoporosis / Secondary Hyperparathyroidism while severe deficiency causes Rickets in children and Osteomalacia in adults. Prevalence of Vitamin D deficiency is approximately >50% specially in the elderly. This assay is useful for diagnosis of vitamin D deficiency and Hypervitaminosis D. It is also used for differential diagnosis of causes of Rickets & Osteomalacia and for monitoring Vitamin D replacement therapy.



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Thyroid Profile Total

Triiodothyronine (T3) <i>ECLIA</i>	0.87	ng/mL	0.57 - 1.78
Total Thyroxine (T4) <i>CLIA</i>	6.7	g/dL	5.1 - 14.1
Thyroid Stimulating Hormone (Ultrasensitive) <i>Chemiluminescence Immuno Assay (CLIA)</i>	3.2	μIU/mL	0.4 - 4.2

Interpretation:

Pregnancy	Reference ranges TSH
1st Trimester	0.1 - 2.5
2nd Trimester	0.2 - 3.0
3rd Trimester	0.3 - 3.0

Note:

TSH levels are subject to circadian variation, reaching peak levels between 2-4 am. and at a minimum between 6-10 pm. The variation is of 50 %, hence time of the day has influence on the measured serum TSH concentrations.

Clinical Use:

- Diagnose Hypothyroidism and Hyperthyroidism
- Monitor T4 replacement or T4 suppressive therapy
- Quantify TSH levels in the subnormal range

Increased Levels : Primary hypothyroidism, Subclinical hypothyroidis, TSH dependent Hyperthyroidism, Thyroid hormone resistance

Decreased Levels: Grace disease, Autonomous thyroid hormone secretion, TSH deficiency

Primary malfunction of the thyroid gland may result in excessive (hyper) or below normal (hypo) release of T3 or T4. In addition as TSH directly affects thyroid function, malfunction of the pituitary or the hypo - thalamus influences the thyroid gland activity. Disease in any portion of the thyroid-pituitary-hypothal- mus system may influence the levels of T3 and T4 in the blood. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels may be low. In addition, in the Euthyroid Sick Syndrome, multiple alterations in serum thyroid function test findings have been recognized in patients with a wide variety of non-thyroidal illnesses (NTI) without evidence of preexisting thyroid or hypothalami c-pituitary diseases. Thyroid Binding Globulin (TBG) concentrations remain relatively constant in healthy individuals. However, pregnancy, excess estrogen's, androgen's, antibiotic steroids and glucocorticoids are known to alter TBG levels and may cause false thyroid values for Total T3 and T4 tests.

TSH	T4	T3	INTERPRETATION
High	Normal	Normal	Mild (subclinical) hypothyroidism
High	Low	Low or Normal	Hypothyroidism
Low	Normal	Normal	Mild (subclinical) hyperthyroidism
Low	High or normal	High or normal	Hyperthyroidism
Low	Low or normal	Low or normal	Nonthyroidal illness; pituitary (secondary) hypothyroidism



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Test Description	Value(s)	Unit(s)	Reference Range
Normal High High Thyroid hormone resistance syndrome (a mutation in the thyroid hormone receptor decreases thyroid hormone function)			

Testosterone Total

Testosterone Total CLIA	354	ng/dL	249 - 836
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Interpretation:

Age in Years	Reference Ranges ng/dL
Males 20-49	249 - 836
Males ≥ 50 years	193 - 740
Females 20-49	8.4 - 48.1
Females ≥ 50	2.9- 40.8

Reference values for Males (7-18 years) characterized by Tanner Stage

Tanner Stage	5-95th percentiles (ng/dL)
1	2.31 - 30.28
2	3.75 - 282.06
3	8.65 - 681.78
4	17.88 - 785.6
5	13.27 - 906.15

Reference values for females (8-18 years) characterized by Tanner Stage

Tanner Stage	5-95th percentiles (ng/dL)
1	0.58 - 33.17
2	4.33 - 23.07
3	6.92 - 42.97
4	15.29 - 1.86
5	15.00 - 102.38

Note

- All applications that require measurement of very low level of testosterone (eg hypogonadal men, children, virilization or intersex disorders in women etc) recommended test is Testosterone total, Ultrasensitive
- LC-MS/MS is the gold standard for steroid hormone assays due to increased sensitivity & specificity as compared to immunoassays

Clinical Use

Assessment of testicular function in males

Increased levels

- Precocious puberty (Males)
- Androgen resistance
- Testotoxicosis
- Congenital Adrenal Hyperplasia



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All Lab results are subject to clinical interpretation by qualified medical professional and this report is not subject to use for any medico-legal purpose.

Patient NAME : Mr DUMMY PL257M	Report STATUS : Final Report		
DOB/Age/Gender : 25 Y/Male	Barcode NO : ZF468880		
Patient ID / UHID : 10573467/OF10573467	Sample Type : Serum		
Referred BY : Self	Report Date : Nov 27, 2024, 01:47 PM.		
Sample Collected : Nov 26, 2024, 01:47 PM			

Test Description	Value(s)	Unit(s)	Reference Range
<p>Decreased levels</p> <ol style="list-style-type: none"> 1. Delayed puberty (Males) 2. Gonadotropin deficiency 3. Testicular defects 4. Systemic diseases 			



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Lipoprotein (A)

Lipoprotein A (Lipo A) <i>Tina-quant</i>	18.0	mg/dL	-
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Interpretation:

Note:
Lp(a) is considered an important risk factor for CHD especially among Indians as Indians tend to have high prevalence of elevated levels of Lp(a)

Lp(a) in mg/dL	REMARKS
(As per Lipid Association of India 2016)	
<30	Low risk
30-49	Moderate Risk
>= 50	High risk

Comments:

Lipoprotein (a) [Lp(a)] consists of an LDL particle that is covalently bound to an additional protein, apolipoprotein (a) [Apo(a)]. Apo(a) has high-sequence homology with the coagulation factor plasminogen and, like LDL, Lp(a) contains apolipoprotein B100 (ApoB). Thus, Lp(a) is both proatherogenic and prothrombotic. Lp(a) is an independent risk factor for Coronary Heart Disease (CHD), Ischemic Stroke, and Aortic Valve Stenosis. Lp(a) is highly heterogenous molecule; the degree of atherogenicity of the Lp(a) particle may depend on the molecular size of the Lp(a)-specific protein. Serum concentrations of Lp(a) are related to genetic factors, and are largely unaffected by diet, exercise and lipid -lowering pharmaceuticals. However, in a patient with additional modifiable CHD risk factors, more aggressive therapy to normalize these factors may be indicated if the Lp(a) value is also increased.

Usage

- Evaluation of increased risk for cardiovascular disease and events:
1. In individuals at intermediate risk for cardiovascular disease
 2. In patients with early atherosclerosis or
 3. In patients with strong family history of early CHD



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Test Description	Value(s)	Unit(s)	Reference Range
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Cortisol, Morning Sample

Cortisol <i>CLIA</i>	13.2	µg/dL	
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Interpretation:

CORTISOL, MORNING	6.7 – 22.60
CORTISOL, EVENING	< 10
CORTISOL, RANDOM	3.0 – 22.6

Note:

Cortisol is best measured in the morning when evaluating for possible Adrenal Insufficiency and best measured in the afternoon or evening to differentiate normal and Cushings Syndrome subjects. Diurnal rhythmicity of cortisol is increased by systemic disease and stress.

Clinical Use :

Direct assessment of Adrenal function

Increased levels: Cushings Syndrome, Ectopic ACTH syndrome, Ectopic CRH syndrome, Adrenal adenoma / carcinoma, Adrenal micronodular dysplasia, Adrenal macronodular hyperplasia, Stress

Decreased Levels: Addisons disease, Pituitary dysfunction



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Apolipoproteins A1 & B

Apolipoprotein A-1 (APO-A) <i>Tina-quant</i>	198	mg/dL	104 - 202
Apolipoprotein B (APO-B) <i>Tina-quant</i>	102	mg/dL	66 - 144
Apo B / Apo A1 Ratio	0.52		0.35 - 0.98

Interpretation:
 The Apo A1 and Apo B blood tests are crucial for assessing lipid metabolism and cardiovascular risk. High Apo A1 and low Apo B levels are associated with a lower risk of cardiovascular disease, while low Apo A1 and high Apo B levels indicate a higher risk. Management involves lifestyle modifications, medications, and regular monitoring to maintain optimal lipid levels and reduce cardiovascular risk. Consulting healthcare providers for accurate interpretation and tailored treatment plans is essential for effective management.

*** End Of Report ***



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