



Patient Name : Mr. BABU ENGANDIYUR Sample UID No. : G4112249F

 Age / Gender
 : 49 Y / Male
 Sample Collected On
 : 24-08-2025 09:46

 Patient ID
 : QLD111700
 Registered On
 : 24-08-2025 17:07

 Referred By
 : DR KEERTHANA RANI
 Reported on
 : 25-08-2025 07:49

Referral Client : CITICARE MEDICAL CENTER(INSURANCE) External Patient ID : 43480 Emirates ID / Passport No : 784197641835414 Print Version : V.1

Department of BIOCHEMISTRY

InvestigationResultsFlagUnitsBiological Reference IntervalMethodGLUCOSE (FASTING)250Hmg/dL74-109Hexokinase

Sample: Fluoride Plasma

Comments:

CLINICAL IMPLICATIONS:

ADA criteria for definitive test for diabetes:

- 1) Fasting blood glucose > 126 mg/dl (> 6.99 mmol/l) on at least two occasions.
- 2) Symptoms of diabetes plus random blood glucose > 200 mg/dl (> 11.1 mmol/l)
- 3) OGTT with 2 hrs. post load (75 gm glucose load) > 200 mg/dl (> 11.1 mmol/l) 4)HbA1c > 6.5%

INTERFERING FACTORS:

- 1) Steroids, diuretics, pregnancy, surgical procedures, anesthesia, obesity, smoking may cause elevated glucose levels.
- 2) Hematocrit > 55%, intense exercise, drug intake may cause lowered glucose level.
- 3)Dawn Phenomenon-Increase in blood glucose typically between 4.00am and 8.00 am due to counter-regulatory hormones.

RECOMMENDATION: As mild borderline cases may present with normal fasting glucose levels, recommended repeat testing

different day.

REFERENCE: 1) Manual of Laboratory and Diagnostics -Frances Fischbach Marshall B. Dunning III [9th Edition]

2) Tietz clinical guide to Laboratory tests(Fourth edition) ALAN H.B.WU

"QLabs compliance with ISO 15189:2022 standards"

Maqsood Rahman Lab Technologist

DHA No:48036476-001



Dr. Vidhya Mohan Specialist Clinical Pathologist Clinical Pathologist DHA No. 23553203-004 Dr. Dheepa Manoharan Medical Director Specialist Microbiologist DHA No. 00231751-004

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LIPID PROFILE TEST

<u>Investigation</u>	<u>Results</u>	Flag	<u>Units</u>	Biological Reference Interval	<u>Method</u>
CHOLESTEROL (TOTAL)	245	Н	mg/dl	Desirable: < 200 Borderline High: 200-239 High: > 240	Enzymatic colorimteric assay
TRIGLYCERIDES	67		mg/dl	Normal Up to 150 Borderline-High 150-199 High 200-499 Very High > 500	Enzymatic colorimetric test
HDL CHOLESTEROL	79		mg/dl	High risk up to 40 Low risk > 60	Homogeneous Enzymatic Colorimetric
LDL CHOLESTEROL DIRECT	145	н	mg/dl	Optimal up to < 100 Near Optimal: 100-129 Borderline: 130-159 High: 160-189 Very High: > 190	Enzymatic, colorimetric method
VLDL CHOLESTEROL	13		mg/dl	10-35	Calculation
NON-HDL CHOLESTEROL	166	Н	mg/dl	Desirable < 130 Borderline 130 – 159 High >159	Calculation
TOTAL CHOLESTEROL / HDL RATIO	3.1			< 4.5	Calculation
LDL / HDL RATIO	1.8			Low Risk < 3.0 Borderline 3.1-6.0 High Risk >6.0	Calculation

Interpretation Notes:

CLINICAL IMPLICATIONS:

- 1. Cholesterol testing evaluates the risk for atherosclerosis, myocardial occlusion, and coronary artery occlusion. Elevated cholesterol levels are a major component in the hereditary hyper lipoproteinemia. It is also used to monitor effectiveness of diet, medications, lifestyle, and stress management.
- 2. The cholesterol to HDL ratio provides more information than does either value alone. When a slightly increased cholesterol is due to high HDL, therapy is not indicated.
- 3. LDL cholesterol has a longer shelf life and determines the CHD risk.

INTERFERING FACTORS:

- 1. Seasonal and positional variations may alter cholesterol levels. Estrogens, ascorbic acid, bilirubin decrease the cholesterol levels
- .Pregnancy, bile salt, high saturated fat, and high cholesterol diet may increase the cholesterol values. Prolonged fasting with ketosis may increase the value.
- 2. Increased levels of HDL may be associated with estrogen therapy, drugs like steroids, alcohol and insulin therapy. Decreased levels are associated with stress, recent illness, starvation, obesity, smoking, hyper triglyceridemia, lack of exercise.
- 3. Increased LDL may be associated with pregnancy, drugs like steroids. Decreased LDL are found in women under estrogen therapy. No fasting may cause false elevation.
- 4. A transient increase in triglycerides occurs following heavy meal, alcohol ingestion, pregnancy, acute illness like cold ,flu, obesity, physical inactivity ,smoking. Transient decrease occurs after strenuous exercise, weight loss.

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Dr. Vidhya Mohan Specialist Clinical Pathologist Clinical Pathologist DHA No. 23553203-004 D. Dl. Dr. Dheepa Manoharan

Medical Director
Specialist Microbiologist
DHA No. 00231751-004

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LIPID PROFILE TEST

<u>Investigation</u> <u>Results</u> <u>Flag</u> <u>Units</u> <u>Biological Reference Interval</u> <u>Method</u> <u>RECOMMENDATION:</u>

- 1. Cholesterol levels >200 mg/dl should be retested and the results averaged and if the results differ by > than 10%, a third test need to be done for confirmation. Perform a comprehensive lipoprotein analysis if cholesterol levels are not lowered within 6 months after start of therapy. If the values are altered in a normal condition, recommended to follow a stable diet for 1 week and overnight fasting (9 to 12 hours) before repeating the test.
- 2. Cholesterol and HDL should not be measured immediately after MI. A 3 month wait is suggested.
- 3. If triglyceride levels are more than 400mg/dl or >10.36mmol/L recommended to fast overnight(9 to 12 hours) and retest .Because of biological and analytical variation, at least 2 serial sample may be necessary for clinical decision making. VLDL cannot be calculated if triglycerides are more than 400mg/dl

REFERENCE: 1) Manual of Laboratory and Diagnostics -Frances Fischbach Marshall B. Dunning III [9th Edition] 2) Tietz clinical guide to Laboratory tests(Fourth edition) ALAN H.B.WU

Sample: Serum

- END OF REPORT -

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