

Patient Name	: Mr. CRISPULO JR GATILA HITALLA	Sample UID No.	: 4091890
Age / Gender	: 49 Y / Male	Sample Collected On	: 30-06-2025 05:00
Patient ID	: QLD069426	Registered On	: 30-06-2025 18:40
Referred By	: CITY CARE	Reported on	: 30-06-2025 22:08
Referral Client	: CITICARE MEDICAL CENTER	External Patient ID	: 46426
Emirates ID / Passport No	:	Print Version	: V.1

**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)**

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
IRON	106		µg/dl	33-193	Ferrozine-no deproteinization

Sample: Serum

Comments :

CLINICAL IMPLICATIONS:

1. The combined results of iron, transferrin, and TIBC are helpful in the differential diagnosis of anaemia ,in assessment of iron deficiency anemia and in the evaluation of thalassemia, sideroblastic anemia and haemochromatosis.
2. Transferrin saturation is a better index of iron saturation. Percent saturation is a better index of iron stores than serum alone. Saturation <15% denotes iron deficiency.

INTERFERING FACTORS:

1. Hemolysis of the blood sample may interfere with testing. Drugs like aspirin, antibiotics, testosterone may cause decreased levels and drugs like ethanol, estrogen may cause an increased iron level.
2. Diurnal variation in iron. Normal values in the morning, low in mid-afternoon, very low in the evening.
3. Serum iron and TIBC may be normal in iron deficiency anemia if Hb is >than 9.0g/dl or >90g/L.

RECOMMENDATION:

In patients receiving folate or Vitamin B12 recommended to repeat iron studies after 1 to 3 months of completion of treatment.

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

- END OF REPORT -

"QLabs compliance with ISO 15189:2022 standards"

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Lab Technologist

DHA No :82488066-001



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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)
LIPID PROFILE TEST

Investigation	Results	Flag	Units	Biological Reference Interval	Method
CHOLESTEROL (TOTAL)	214	H	mg/dl	Desirable: < 200 Borderline High: 200-239 High: > 240	Enzymatic colorimetric assay
TRIGLYCERIDES	358	H	mg/dl	Normal Up to 150 Borderline-High 150-199 High 200-499 Very High > 500	Enzymatic colorimetric test
HDL CHOLESTEROL	53		mg/dl	High risk up to 40 Low risk > 60	Homogeneous Enzymatic Colorimetric
LDL CHOLESTEROL DIRECT	99		mg/dl	Optimal up to < 100 Near Optimal: 100-129 Borderline : 130-159 High: 160-189 Very High: > 190	Enzymatic, colorimetric method
VLDL CHOLESTEROL	72	H	mg/dl	10-35	Calculation
NON-HDL CHOLESTEROL	161	H	mg/dl	Desirable < 130 Borderline 130 – 159 High >159	Calculation
TOTAL CHOLESTEROL / HDL RATIO	4.0	--	--	< 4.5	Calculation
LDL / HDL RATIO	1.9	--	--	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

- END OF REPORT -

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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)
LIPID PROFILE TEST

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
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inactivity ,smoking. Transient decrease occurs after strenuous exercise, weight loss.

RECOMMENDATION:

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

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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)
Renal Function Test

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
UREA (SERUM)	44.3		mg/dl	16.6-48.5	Kinetic test,Urease
CREATININE (SERUM)	1.01		mg/dl	0.7 - 1.2	Alkaline picrate
URIC ACID (SERUM)	6.11		mg/dl	3.4-7	Enzymatic colorimetric test
BLOOD UREA NITROGEN (SERUM)	20.7	H	mg/dl	6-20	Calculation
BUN/CREATININE RATIO	20.50		mg/dl	10-30	Calculation

Interpretation Notes :

CLINICAL IMPLICATIONS:

1. A markedly increased BUN is conclusive of severe impaired glomerular function and in chronic renal disease BUN level correlates better with the symptoms of uremia than does the serum creatinine.
2. Uric acid levels is used most in the evaluation of renal failure, gout, and leukaemia. In gout the amount of increase is not directly related to the severity of the disease. Acute levels may occur following administration of cytotoxic drugs.
3. In chronic renal disease, BUN/creatinine ratio is a better indicator to evaluate the renal problem than evaluating either alone. For each 50% reduction in GFR serum creatinine doubles. In chronic renal disease the plasma levels of creatinine may be more sensitive to changes in glomerular function than creatinine clearance, which may be factitiously higher than the true value.

INTERFERING FACTORS:

1. A combination of low protein high carbohydrate diet, late pregnancy (PHYSIOLOGIC HYDREMIA), IV feedings may cause a low level of BUN. BUN is normally lower in children and women than adult. Elderly person can have an increased BUN level. Many drugs like steroids, tetracyclines, thyroxine, stress, strenuous exercise may alter the values of BUN levels.
2. Stress, strenuous exercise, purine rich diet(liver, kidney, sweet breads)increases uric acid levels. High levels of aspirin, low purine intake, coffee, tea intake may cause a decreased level of uric acid. Drugs like steroids, diuretics, acetaminophen may alter the uric acid levels.
3. High levels of ascorbic acid, cephalosporin, diet high in meat, ketoacidosis may increase serum creatinine substantially. Creatinine is falsely decreased by bilirubin, glucose, histamine, quinidine compounds. Drugs like cephalosporins may alter the values. Lipemic and homolyzed samples may cause false elevations.

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

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**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)**

GLYCATED HEMOGLOBIN (Hba1c)

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
Sample: EDTA Whole Blood					
GLYCATED HEMOGLOBIN (HbA1C) ^	5.9		%	<5.7 non-diabetic 5.7-6.4 Pre-Diabetic >6.4 Diabetic	Turbidimetric Inhibition Immunoassay
Average Blood Glucose	122.6		mg/dl	90-120 Good Control 121-150 Fair Control 151-180 Unsatisfactory Control >180 Poor Control	Calculated

Comments :

False elevated levels may be due to hypertriglyceridemia, iron deficiency anemia, B12 deficiency, vit C supplement usage, uremia, hemoglobinopathies. In such cases recommended to evaluate the value using alternative index like fructosamine, glycated albumin or continuous glucose monitoring

CLINICAL IMPLICATIONS:

- 1) Glycated hemoglobin reflects average blood sugar level for 2 to 3 month period and useful for evaluating diabetic medications and to track the control of blood glucose in milder cases.
- 2) Increase in Glycated hemoglobin occurs in non diabetic conditions like Iron deficiency anemia, splenectomy, alcohol toxicity. Decrease in Glycated Hemoglobin in hemolytic anemia, chronic blood loss, pregnancy and chronic renal failure.
- 3) Improvement in the glucose control occurring in the 4 weeks before drawing of the sample is not well reflected in the result since the formation of glycated haemoglobin is irreversible.

INTERFERING FACTORS:

- 1) Presence of HbF and HbH cause falsely elevated values.
- 2) Presence of Hb S, C, E, D, G and Lepore cause falsely decrease results.
- 3) If test results are not consistent with clinical finding check the patient for HbF which elevates HbA1c results.
- 4) Haemolytic blood samples may cause falsely low results because of increased erythrocyte turnover.

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

- END OF REPORT -

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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)
LIVER FUNCTION TEST

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
ALT / SGPT	26.3		U/L	10-50	IFCC with P5P
AST / SGOT	28		U/L	10-50	IFCC with P5P
ALP (ALKALINE PHOSPHATASE)	84		U/L	40-129	Colorimetric assay
GGT (GAMMA GLUTAMYL TRANSFERASE)	55		U/L	8-61	Enzymatic colorimetric assay
BILIRUBIN (TOTAL)	0.2		mg/dl	0.1-1.2	Diazo
BILIRUBIN (DIRECT)	0.08		mg/dl	0-0.3	Diazo
INDIRECT BILIRUBIN	0.12		mg/dl	0.00-1.1	Calculated Parameter
TOTAL PROTEIN	8.2		g/dl	6.6-8.7	Colorimetric assay
ALBUMIN (SERUM)	4.6		g/dl	3.97-4.94	Colorimetric assay
GLOBULIN	3.6	H	g/dl	2.35 - 3.5	Calculated Parameter
A/G RATIO	1.3			1.1-2.5	Calculated Parameter

Interpretation Notes :

CLINICAL IMPLICATIONS:

- 1) Total Bilirubin elevation accompanied by jaundice is due to hepatic, obstructive, hemolytic and blood group compatibility.
- 2) Increase albumin is associated with dehydration and decrease is due to acute and chronic inflammation, burns and heart failure.
- 3) Although AST levels always increase in acute MI, ALT level doesn't always increase unless there also liver damage.
- 4) ALT is usually increased more than AST in acute extra hepatic biliary obstruction.
- 5) ALT is more specific than AST for liver disease but AST is more sensitive to liver disease.
- 6) Alkaline phosphatase normal values are higher in pediatric patient and in pregnancy. Values may increase up to 3 times in puberty.
- 7) GGT is used to confirm biliary abnormality and is elevated in hepatobiliary disease but not in uncomplicated bone disease.

- END OF REPORT -

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**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)
LIVER FUNCTION TEST**

Investigation	Results	Flag	Units	Biological Reference Interval	Method
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8) GGT values are higher in new born, first 3 to 6 month . Adult male have 25% higher values than female.

INTERFERING FACTORS:

- 1) Certain foods like carrots, yam, drugs, anorexia, prolonged fasting may falsely increase bilirubin level.
- 2) Albumin levels may reduce in pregnancy, over hydration, edema, drugs, obesity.
- 3) Young children, pregnant women, post-menopausal women have physiological high level of ALT. Alkaline phosphatase increase after fatty meal.
- 4) Slight reduce level of AST can be seen during pregnancy and false reduced level in severe liver disease.

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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Department of HEMATOLOGY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)
ADVANCE COMPLETE BLOOD COUNT

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
HEMOGLOBIN	14.3		g/dl	13-17	photometric
WBC COUNT	8		10 ³ /uL	4-11	Electrical Impedance
RBC COUNT	5.05		10 ⁶ /uL	4.5-5.5	Electrical Impedance
HEMATOCRIT	43		%	42-52	Calculation
MCV	85		fL	78-100	Calculation
MCH	28.3		pg	27-31	Calculation
MCHC	33.3		g/dl	31-35	Calculation
RDW	15.3		%	9.3-16	Calculation
RDW-SD	46.8		fL	38.9-49	Calculation
MPV	8.3	L	fL	8.8-12.5	Calculation
PLATELET COUNT	296		10 ³ /uL	150-400	Electrical Impedance
* Neutrophil	55.24		%	40-80	VCS-Method
* Lymphocyte	32.47		%	20-40	VCS-Method
* Monocyte	6.11		%	2-10	VCS-Method
* Eosinophil	4.99		%	1-8	VCS-Method
* Basophil	1.19		%	0-2	VCS-Method
* ABSOLUTE NEUTROPHIL COUNT	4.44		10 ³ /uL	1.5-7	Calculation
* ABSOLUTE LYMPHOCYTE COUNT	2.61		10 ³ /uL	1.5-4	Calculation
* ABSOLUTE MONOCYTE COUNT	0.49		10 ³ /uL	0-0.8	Calculation
* ABSOLUTE EOSINOPHIL COUNT	0.4		10 ³ /uL	0-0.6	Calculation
* ABSOLUTE BASOPHIL COUNT	0.1		10 ³ /uL	0-0.2	Calculation

Interpretation Notes :

Interfering factors: Factors such as age, gender, pregnancy, drug intake, excessive fluid intake, dehydration, hyperlipidemia, stress, exercise, post-operative state, new born, clotted specimen may interfere with test results. Hence recommended fresh EDTA blood sample for confirmation.

Reference :-Manual of Laboratory and Diagnostics -Frances Fischbach Marshall B. Dunning III [9th Edition

Sample: EDTA Whole Blood

- END OF REPORT -

Note:

"The analytes with asterix (*) symbol are non-accredited parameters."
"QLabs compliance with ISO 15189:2022 standards"



Ebin C Lorance
Lab Technologist



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**Department of IMMUNOLOGY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)**

Investigation	Results	Flag	Units	Biological Reference Interval	Method
VITAMIN D, 25-OH (TOTAL)	30.6		ng/mL	Deficient : ≤ 20 insufficient: 21-29 Sufficient: ≥ 30 Toxicity : >80	ECLIA

Sample: Serum

Comments :

CLINICAL IMPLICATIONS:

1. Increased Vitamin D levels are seen in gastrointestinal symptoms like anorexia, nausea, vomiting, constipation, hypercalcemia, renal colic, supplements, normal growing children, pregnant and lactating females, tuberculosis, idiopathic hypercalciuria, sarcoidosis. Levels can increase to 200 -300pg/ml during treatment of osteoma Lacia with physiological doses of vitamin D.
2. Decreased levels are seen in Inadequate diet, Inadequate exposure to sunlight, liver disease, Malabsorption syndrome, osteoma Lacia, Anticonvulsants, rickets, chronic renal failure, pseudohypoparathyroidism, post-menopausal osteoporosis and adults with insulin requiring diabetes mellitus.
3. 25(OH) levels do not indicate clinical vitamin D status in patients with chronic renal failure or type 1 vitamin D dependent rickets or when calcitriol is used as a supplement.

INTERFERING FACTORS:

Age, season of the year, diarrhoea or vomiting, certain drugs, diseases, and long term hyperalimentation are the factors that may interfere with the vitamin levels.

RECOMMENDATION:

Recommended to evaluate alternate cause of impaired mineralization ,if the levels are not consistent with the suspected diagnosis.

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
- 3) Clinical microbiology procedures 4th edition AMY L LEBER

- END OF REPORT -

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**Department of IMMUNOLOGY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)**

Investigation	Results	Flag	Units	Biological Reference Interval	Method
THYROID STIMULATING HORMONE (TSH)	2.57		uIU/mL	0.27-4.2	ECLIA

Sample: Serum

Comments :

CLINICAL IMPLICATIONS:

- 1.TSH has diurnal rhythm, peaks at 2:00-4:00am and has low levels at 5:00-6:00pm with ultradian rhythm (shorter than circadian).
2. Moderately high TSH is often found in euthyroid patients during treatment for hyperthyroidism. In treated hyperthyroid patient, TSH may remain low for 4-6 week after euthyroid state is achieved. TSH surges with birth,peaking at 30min at 25-160mU/L,declining to cord blood levels by 3 days,and reaching adult values in the first week of life.
3. If there is clear evidence of hypothyroidism and TSH is not elevated then an implication of possible hypopituitarism exists.

INTERFERING FACTORS:

- 1.Values are normally high in neonatal cord blood,and comes to normal by first week.
- 2.Values are increased in elderly patients,drugs like amphetamine abuse,potassium iodide,lithium,iodine containing drugs.
- 3.Values may be decreased in first trimester of pregnancy and during treatment with thyroxine and corticosteroids.
- 4.Heterophilic antibodies may falsely decrease or increase test results.

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

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Patient Name	: Mr. CRISPULO JR GATILA HITALLA	Sample UID No.	: 4091890
Age / Gender	: 49 Y / Male	Sample Collected On	: 30-06-2025 05:00
Patient ID	: QLD069426	Registered On	: 30-06-2025 18:40
Referred By	: CITY CARE	Reported on	: 30-06-2025 22:08
Referral Client	: CITICARE MEDICAL CENTER	External Patient ID	: 46426
Emirates ID / Passport No	:	Print Version	: V.1

**Department of IMMUNOLOGY
GENERAL WELLNESS CHECK-UP LITE(CITICARE)**

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
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- END OF REPORT -

"QLabs compliance with ISO 15189:2022 standards"

Mohamed Babiker

Mojahid Babiker
Lab Technologist

DHA No :82488066-001



Dr. Vidhya Mohan

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Dr. Dheepa Manoharan

Dr. Dheepa Manoharan
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