

Patient Name	: Ms. ANNA DANILINA	Sample UID No.	: 4071531
Age / Gender	: 44 Y / Female	Sample Collected On	: 16-04-2025 13:28
Patient ID	: QLD071405	Registered On	: 16-04-2025 13:39
Referred By	: PESHAWAR	Reported on	: 16-04-2025 21:17
Referral Client	: CITICARE MEDICAL CENTER	External Patient ID	: 46509
Emirates ID / Passport No	:	Print Version	: V.1

Department of BIOCHEMISTRY

Investigation	Results	Flag	Units	Biological Reference Interval	Method
HOMOCYSTEINE	8.7		umol/L	Refer interpretation notes.	Enzymatic Assay, Novel Enzyme Cycling assay

Sample: Serum

Comments :

Age, pregnancy and renal function are important. The intake of folic acid as either supplements or through fortification of foods must also be considered.

Group	Folate supplemented	Non supplemented
Pregnancy	< 8	< 10
Children < 15 Years	< 8	< 10
Adult 15 - 65 Years	< 12	< 15
Elderly > 65	< 16	< 20

CLINICAL IMPLICATIONS:

1. Increased or elevated levels of homocysteine (Hcys) are found in Folic acid deficiency, Vitamin B12 deficiency, Homocystinuria. Serum or plasma Hcys is used as a functional test for Vitamin B12. folate, Vitamin B6 deficiency. Elevated Hcys is also an independent risk factor for cardiovascular disease.
2. Intraindividual variability is 8% over a 1 year period, although it can be as much as 25% in patients with hyperhomocysteinemia.

INTERFERING FACTORS:

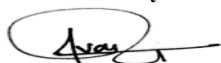
1. Serum results are 5-10% higher than EDTA or heparin plasma, whereas citrated plasma results are 5-15% lower. Hcys increase 10-15% increase 6-8 hr after a high protein meal.
2. Penicillamine reduces plasma levels of Hcys. Nitrous oxide, methotrexate deficiency and azauridine increase plasma levels of homocysteine.

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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Verified By



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DHA No. 87250933-002



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Department of IMMUNOLOGY

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
FERRITIN	128	H	µg/L	10-120 Iron overload : > 200	ECLIA

Sample: Serum

Comments :

CLINICAL IMPLICATIONS:

1. Ferritin is the most reliable indicator of total body iron status and is more specific and sensitive than the iron concentration or TIBC for diagnosing iron deficiency. Ferritin decreases before anaemia and other changes occurs.
2. Decreased ferritin usually indicates iron deficiency anaemia and increased ferritin occurs in iron excess and other causes like hyperthyroid, liverdisease, haemolytic, megaloblastic, sideroblastic and thalassemia.
3. Ferritin levels less than 10ng/ml usually indicate iron deficiency anaemia. In conditions of iron overload and in some chronic diseases, serum ferritin is an unreliable estimate of storage iron and in such cases serum ferritin is less sensitive than serum iron concentration, TIBC, or percent transferrin saturation.

INTERFERING FACTORS:

1. Recently administered radioactive medication can cause spurious results.
2. Hemolyzed blood, age, diet, antithyroid therapies are the factors that may interfere the ferritin values.
3. Ferritin is not of value to evaluate iron stores in alcoholic person with 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
- 3) Clinical microbiology procedures 4th edition AMY L LEBER

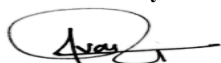
RECOMMENDATION:

Recommended to do Iron, Ferritin, TIBC together to distinguish between iron deficiency anaemia and the anaemia of chronic disease.

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Department of ENDOCRINOLOGY

Investigation	Results	Flag	Units	Biological Reference Interval	Method
* THYROID PEROXIDASE (MICROSOMAL) ANTIBODY (A-TPO)	< 9		IU/mL	< / =34	ECLIA

Sample: Serum

Comments :

Clinical Implication: 1) High titers of thyroglobulin and thyroid microsomal antibodies (>1:400) are found in Hashimoto's disease, but elevation can also be seen in other autoimmune diseases. 2) Increased thyroid antibodies also occur in Graves' disease; thyroid carcinoma; idiopathic myxedema, pernicious anemia; SLE, RA Sjogren's syndrome; subacute thyroiditis; nontoxic nodular goiter.

Interfering factors: 1) About 10% of the normal population may have low levels of thyroid antibodies with no symptoms of the disease. Incidence of low titer is higher in women and increases with age. 2) Antibody production may be confined to lymphocytes within the thyroid, resulting in negative serum test results.

Reference <https://emedicine.medscape.com>

Interpretation Notes :

Anti-thyroid peroxidase (anti-TPO) antibodies are specific against TPO, which catalyses iodine oxidation & iodination reactions in the thyroid gland.

10-15% of normal individuals & during pregnancy can have raised anti-TPO antibody titers.

Anti-TPO antibodies are the most common anti-thyroid autoantibody, present in approximately 90% of Hashimoto's thyroiditis, 75% of Graves' disease, and 10-20% of nodular goiter or thyroid carcinoma. Relatives of patients with an autoimmune thyroid disorder (40%-50%) may have elevated serum TPOAb levels. High serum antibodies are found in active phase chronic autoimmune thyroiditis & antibody titer can be used to assess disease activity. Other autoimmune disorders often associated with elevated TPOAb include Sjogren's syndrome, lupus, rheumatoid arthritis, diabetes mellitus (type I), and pernicious anemia.

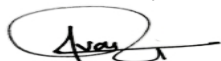
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Note:

"The analytes with asterisk (*) symbol are non-accredited parameters."

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Department of HAEMATOLOGY

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
ERYTHROCYTE SEDIMENTATION RATE (ESR)	08		mm/hr	0-20	Automated

Sample: EDTA Whole Blood

Comments :

CLINICAL IMPLICATIONS:

1.ESR is is not diagnostic of any particular disease ,but rather is an indication that the disease process is ongoing and must be investigated.It is also useful in monitoring the progression of inflammatory disease.

2.ESR is most useful in the diagnosis of temporal arteritis,rheumatoid arthritis and polymyalgia rheumatica.If the patient is treated with steroids,the ESR will decrease with clinical improvement.ESR should not be used to screen asymptomatic patient for disease .

INTERFERING FACTORS:

1.Anemia,young children,menstruation,macrocytosis,,presence of fibrinogen,globulins,C-reactive protein,high cholesterol,pregnancy after 12 weeks until about the fourth postpartum week,certain drugs like heparin can cause an increased ESR levels.ESR may be very high (upto 60 mm /hr)in apparently healthy woman aged 70 to 89 years.

2.Factors like high blood sugar,high albumin,high phospholipids,decreased fibrinogen level in the blood in newborn,hypofibrinogenemia,High Hb and RBC,Polycythemia,High WBC,Abnormal RBC(sickle cells,spherocytes,microcytosis,)high WBC,certain drugs like steroids,aspirin causes reduced ESR levels.

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**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)**

Investigation	Results	Flag	Units	Biological Reference Interval	Method
IRON	134		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.

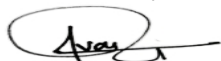
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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)
LIPID PROFILE TEST

Investigation	Results	Flag	Units	Biological Reference Interval	Method
CHOLESTEROL (TOTAL)	159		mg/dl	Desirable: < 200 Borderline High: 200-239 High: > 240	Enzymatic colorimetric assay
TRIGLYCERIDES	59		mg/dl	Normal Up to 150 Borderline-High 150-199 High 200-499 Very High > 500	Enzymatic colorimetric test
HDL CHOLESTEROL	66		mg/dl	High risk up to 40 Low risk > 60	Homogeneous Enzymatic Colorimetric
LDL CHOLESTEROL DIRECT	84		mg/dl	Optimal up to < 100 Near Optimal: 100-129 Borderline : 130-159 High: 160-189 Very High: > 190	Enzymatic, colorimetric method
VLDL CHOLESTEROL	12		mg/dl	10-35	Calculation
NON-HDL CHOLESTEROL	93		mg/dl	Desirable < 130 Borderline 130 – 159 High >159	Calculation
TOTAL CHOLESTEROL / HDL RATIO	2.4		--	< 4.5	Calculation
LDL / HDL RATIO	1.3	L	--	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.

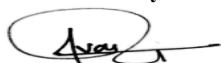
INTERFERING FACTORS:

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

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**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)
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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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.

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

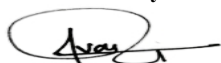
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Sample: Serum

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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)
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)	19.2		mg/dl	16.6-48.5	Kinetic test,Urease
CREATININE (SERUM)	0.57		mg/dl	0.5 - 0.9	Alkaline picrate
URIC ACID (SERUM)	3.48		mg/dl	2.4-5.7	Enzymatic colorimetric test
BLOOD UREA NITROGEN (SERUM)	9.0		mg/dl	6-20	Calculation
BUN/CREATININE RATIO	15.79		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:

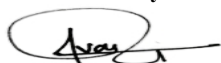
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Sample: Serum

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**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)**

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) ^	4.8		%	<5.7 non-diabetic 5.7-6.4 Pre-Diabetic >6.4 Diabetic	Turbidimetric Inhibition Immunoassay
Average Blood Glucose	91.1		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.

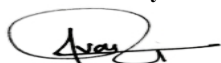
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Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)
LIVER FUNCTION TEST

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
ALT / SGPT	8.74	L	U/L	10-35	IFCC with P5P
AST / SGOT	24		U/L	10-35	IFCC with P5P
ALP (ALKALINE PHOSPHATASE)	59		U/L	35-104	Colorimetric assay
GGT (GAMMA GLUTAMYL TRANSFERASE)	13		U/L	5-36	Enzymatic colorimetric assay
BILIRUBIN (TOTAL)	0.6		mg/dl	0.1-1.2	Diazo
BILIRUBIN (DIRECT)	0.32	H	mg/dl	0-0.3	Diazo
INDIRECT BILIRUBIN	0.28		mg/dl	0.00-1.1	Calculated Parameter
TOTAL PROTEIN	6.7		g/dl	6.6-8.7	Colorimetric assay
ALBUMIN (SERUM)	4.3		g/dl	3.97-4.94	Colorimetric assay
GLOBULIN	2.4		g/dl	2.35 - 3.5	Calculated Parameter
A/G RATIO	1.8			1.1-2.5	Calculated Parameter

Interpretation Notes :

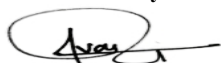
CLINICAL IMPLICATIONS:

- 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 -

"QLabs compliance with ISO 15189:2022 standards"

Verified By



Vaishnav Jayamohan
Lab Technologist

DHA No. 87250933-002



Authorised By



Dr. Dheepa Manoharan
Medical Director
Specialist Microbiologist
DHA No. 00231751-004

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Patient Name	: Ms. ANNA DANILINA	Sample UID No.	: 4071531
Age / Gender	: 44 Y / Female	Sample Collected On	: 16-04-2025 13:28
Patient ID	: QLD071405	Registered On	: 16-04-2025 13:39
Referred By	: PESHAWAR	Reported on	: 16-04-2025 21:17
Referral Client	: CITICARE MEDICAL CENTER	External Patient ID	: 46509
Emirates ID / Passport No	:	Print Version	: V.1

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

Investigation **Results** **Flag** **Units** **Biological Reference Interval** **Method**

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.

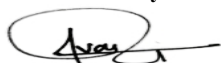
REFERENCE: 1) Manual of Laboratory and Diagnostics -Frances Fischbach Marshall B. Dunning III [9th Edition]
2) Tietz clinical guide to Laboratory tests(Fourth edition) ALAN H.B.WU

Sample: Serum

- END OF REPORT -

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Patient Name	: Ms. ANNA DANILINA	Sample UID No.	: G4071531
Age / Gender	: 44 Y / Female	Sample Collected On	: 16-04-2025 13:28
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**Department of BIOCHEMISTRY
GENERAL WELLNESS CHECK-UP(PESHAWAR)**

GLUCOSE R/F

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
Sample: Fluoride Plasma GLUCOSE FASTING	89		mg/dl	74-109	Hexokinase

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 on a different day.

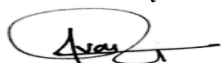
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Patient Name : Ms. ANNA DANILINA
Age / Gender : 44 Y / Female
Patient ID : QLD071405
Referred By : PESHAWAR
Referral Client : CITICARE MEDICAL CENTER
Emirates ID / Passport No :

Sample UID No. : EB4071531
Sample Collected On : 16-04-2025 13:28
Registered On : 16-04-2025 13:39
Reported on : 16-04-2025 21:17
External Patient ID : 46509
Print Version : V.1

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

Investigation	Results	Flag	Units	Biological Reference Interval	Method
HEMOGLOBIN	14.4		g/dl	12-15	photometric
WBC COUNT	5.8		10 ³ /uL	4-11	Electrical Impedance
RBC COUNT	4.28		10 ⁶ /uL	3.8-4.8	Electrical Impedance
HEMATOCRIT	42.8		%	37-47	Calculation
MCV	100.1	H	fL	78-100	Calculation
MCH	33.8	H	pg	27-31	Calculation
MCHC	33.7		g/dl	31-35	Calculation
RDW	11.9		%	9.3-16	Calculation
RDW-SD	41.6		fL	38.9-49	Calculation
MPV	10.9		fL	8.8-12.5	Calculation
PLATELET COUNT	229		10 ³ /uL	150-400	Electrical Impedance
* Neutrophil	47.3		%	40-80	VCS-Method
* Lymphocyte	34.2		%	20-40	VCS-Method
* Monocyte	15.1	H	%	2-10	VCS-Method
* Eosinophil	2.9		%	1-8	VCS-Method
* Basophil	0.5		%	0-2	VCS-Method
* ABSOLUTE NEUTROPHIL COUNT	2.7		10 ³ /uL	1.5-7	Calculation
* ABSOLUTE LYMPHOCYTE COUNT	2		10 ³ /uL	1.5-4	Calculation
* ABSOLUTE MONOCYTE COUNT	0.9	H	10 ³ /uL	0-0.8	Calculation
* ABSOLUTE EOSINOPHIL COUNT	0.2		10 ³ /uL	0-0.6	Calculation
* ABSOLUTE BASOPHIL COUNT	0		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."
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Patient Name	: Ms. ANNA DANILINA	Sample UID No.	: 4071531
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**Department of IMMUNOLOGY
GENERAL WELLNESS CHECK-UP(PESHAWAR)**

Investigation	Results	Flag	Units	Biological Reference Interval	Method
VITAMIN D, 25-OH (TOTAL)	33.3		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.

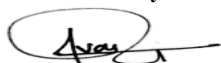
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- 3) Clinical microbiology procedures 4th edition AMY L LEBER

- END OF REPORT -

"QLabs compliance with ISO 15189:2022 standards"

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**Department of IMMUNOLOGY
GENERAL WELLNESS CHECK-UP(PESHAWAR)**

<u>Investigation</u>	<u>Results</u>	<u>Flag</u>	<u>Units</u>	<u>Biological Reference Interval</u>	<u>Method</u>
VITAMIN B12	665		pg/mL	197-771	ECLIA

Sample: Serum

Comments :

CLINICAL IMPLICATIONS: 1. Levels of Vitamin B12 and folate are usually tested in conjunction with one another for the diagnosis of Macrocytic anaemia and measurement of unsaturated VB12 binding capacity is valuable in distinguishing between untreated polycythemia vera and other conditions in which there is an elevated HCT. 2. Serum levels can be low in the absence of either anaemia or macrocytosis(eg. in patients with myeloma, aplastic anaemia) and conversely elevated Transcobalamin II can cause a normal or increase Vitamin B12 levels despite deficient liverstores.

INTERFERING FACTORS: 1. Blood transfusion, pregnancy, elderly patients, high vitamin C and A, smoking, drugs like aminoglycosides, metformin may alter the vitamin B 12 levels. 2. Low serum vitamin B 12 levels often occur in folate deficiency, and B12 deficiency can be masked by folate therapy.

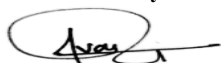
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Department of IMMUNOLOGY
GENERAL WELLNESS CHECK-UP(PESHAWAR)
THYROID FUNCTION TEST (T3,T4,TSH)

Investigation	Results	Flag	Units	Biological Reference Interval	Method
TRIIODOTHYRONINE, TOTAL (T3)	1.05		ng/mL	0.8-2	ECLIA
THYROXINE, TOTAL (T4)	6.4		ug/dL	5.1-14.1	ECLIA
THYROID STIMULATING HORMONE (TSH)	7.32	H	uIU/mL	0.27-4.2	ECLIA

Interpretation Notes :

Total T3 Clinical implication:

Total T3 level is a quantitative determination of the total T3 concentration in the blood and is the test of choice in the diagnosis of T3 thyrotoxicosis.

Total T4 Clinical Implications:

Total T4 is a good index of thyroid function when TB G (Thyroid Binding Globulin) is normal. The increase in TBG levels normally seen in pregnancy and with estrogen therapy increases total T4 levels. The decrease of TBG levels in person receiving anabolic steroids, in chronic liver disease and in nephrosis decrease the total T4 value..

TSH CLINICAL IMPLICATIONS:

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).

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.

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

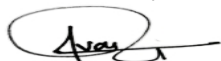
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Sample: Serum

- END OF REPORT -

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