



Name : Mansoor Ali Khan Jaleel Sample Date : 04/01/2023 15:31 PM

Lab. No. : 3023123934 Report Date : 04/01/2023 16:42 PM

Contract. : Nas (WN, VN)-Dubai Branch Doctor references : Dr. Marvis Enyl

Patient No. : 30-86893

File No. : 29239

Branch : Al Borg Lab. Dubai Age : 37 Year DOB 05/12/1986 Sex : Male

Deirah

Cholesterol/HDL/LDL/Triglycerides

Test	Result		Unit	Ref. Range
Cholesterol	203	Н	mg/dl	No risk <200 Moderate risk 200 - 240 High risk >240
Comments Primary sample: Serum Methodology: Enzymatic colorimetric assay in the presence pf peroxidase.				
Triglycerides (TG) in Serum	65		mg/dL	Normal<200
Comments Primary sample: Serum Methodology: Enzymatic colorimetric assay				
HDL Cholesterol	42	L	mg/dL	Optimum level ≥ 60 Borderline risk: 40-59 High risk: <40
Comments Primary sample: Serum Methodology: HOmogeneous Enzymatic colorimetric assay.				
LDL Cholesterol	148	Н	mg/dl	Optimal <100 Near optimal 100-129 Borderline high 130-159 High 160-189 Very high >189

Comments

Sample type: Serum Methodology: Calculation

*This is non-accredited test.

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Cholesterol/HDL/LDL/Triglycerides

Test	Result		Unit	Ref. Range
Non-HDL Cholesterol	161	Н	mg/dL	very high >220 High 190-219 Borderline high 160-189 Near Ideal 130-159 Ideal for people at risk of heart disease <130 Ideal for people at very high

Comments

Patients with elevated levels of non-HDL-C and normal levels of LDL-C, often have an increased number of LDL particles, increased apo B or increased small, dense LDL particles and these are associated with an increased risk of CVD. Therefore non-HDL-C might be more valuable indicator of cardiovascular risk than LDL-C.

The treatment goal for non-HDL cholesterol in persons with high triglycerides (>199 mg/dl) is 30 mg/dl higher than their LDL cholesterol goal.

Reviewed By:

Dr. Shailendra Rathod.MD **Laboratory Director Physician Specialist Clinical Pathology**

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Verified By: Shailendra Rathod.MD

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Age: 37 Year

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Sex

: Male

Chemistry Unit

Test Result Unit Ref. Range 70 - 200 Glucose in Plasma (Random) 100 mg/dl

Comments

Branch

Primary sample: Plasma

Methodology: Enzymatic reference method with hexokinase

**Please note new reference range

Reviewed By:

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Pathology

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Complete Blood Count - (CBC)

Test	Result		Unit	Ref. Range
Hemoglobin	14.6		g/dL	13.5 - 17.5
Hematocrit	43.3		%	38.8 - 50
Red cell count	5.09	;	x10¹²/L	4.32 - 5.72
MCV	85.1		fL	81.2 - 95.1
MCH	28.7		pg	26.5 - 32.6
MCHC	33.7		g/dL	32 - 36
RDW	13.0		%	11.8 - 15.6
Total Leucocytic Count	7.66	:	x10º/L	3.5 - 10.5
Basophils absolute count	0.03	:	x10°/L	0 - 0.3
Basophils relative count	0.39		%	
Eosinophils absolute count	0.15	:	x10°/L	0.05 - 0.5
Eosinophils relative count	1.96		%	
Neutrophils absolute count	3.20	;	x10º/L	1.7 - 7
Neutrophils relative count	41.78		%	
Lymphocytes absolute count	3.71	Н	x10º/L	0.9 - 3.1
Lymphocytes relative count	48.43		%	
Monocytes absolute count	0.57	;	x10º/L	0.3 - 0.9
Monocytes relative count	7.44		%	

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Deirah

Complete Blood Count - (CBC)

 Test
 Result
 Unit
 Ref. Range

 Platelet Count
 308
 x10°/L
 150 - 450

Comments

Sample type: EDTA Whole Blood

Methodology: (Hemoglobin = Cyanmethemoglobin) (Hematocrit = Calculation) (RBC, PLT, WBC and Diff Count = Flow cytometry)

Reviewed By:

Dr. Shailendra Rathod.MD

Laboratory Director

Physician Specialist Clinical

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