

Patient Information	Specimen Information	Client Information

COMMENTS: FASTING: YES

Test Name	In Range	Out Of Range	Reference Range	Lab
THYROID PANEL WITH TSH				
THYROID PANEL				TP
T3 UPTAKE	29		22-35 %	
T4 (THYROXINE), TOTAL	7.8		5.1-11.9 mcg/dL	
FREE T4 INDEX (T7)	2.3		1.4-3.8	
TSH	3.38		mIU/L	TP
			Reference Range	
			> or = 20 Years 0.40-4.50	
			Pregnancy Ranges	
			First trimester 0.26-2.66	
			Second trimester 0.55-2.73	
			Third trimester 0.43-2.91	
LIPID PANEL, STANDARD				
CHOLESTEROL, TOTAL	173		<200 mg/dL	TP
HDL CHOLESTEROL	58		> OR = 50 mg/dL	TP
TRIGLYCERIDES	59		<150 mg/dL	TP
LDL-CHOLESTEROL		101 H	mg/dL (calc)	TP
Reference range: <100				
Desirable range <100 mg/dL for primary prevention; <70 mg/dL for patients with CHD or diabetic patients with > or = 2 CHD risk factors.				
LDL-C is now calculated using the Martin-Hopkins calculation, which is a validated novel method providing better accuracy than the Friedewald equation in the estimation of LDL-C. Martin SS et al. JAMA. 2013;310(19): 2061-2068 (http://education.QuestDiagnostics.com/faq/FAQ164)				
CHOL/HDL-C RATIO	3.0		<5.0 (calc)	TP
NON HDL CHOLESTEROL	115		<130 mg/dL (calc)	TP
For patients with diabetes plus 1 major ASCVD risk factor, treating to a non-HDL-C goal of <100 mg/dL (LDL-C of <70 mg/dL) is considered a therapeutic option.				
COMPREHENSIVE METABOLIC PANEL				TP
GLUCOSE	89		65-99 mg/dL	
			Fasting reference interval	
UREA NITROGEN (BUN)	17		7-25 mg/dL	
CREATININE	0.76		0.50-1.03 mg/dL	
EGFR	95		> OR = 60 mL/min/1.73m ²	
BUN/CREATININE RATIO	SEE NOTE:		6-22 (calc)	
	Not Reported: BUN and Creatinine are within reference range.			
SODIUM	140		135-146 mmol/L	
POTASSIUM	4.6		3.5-5.3 mmol/L	

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CHLORIDE	107		98-110 mmol/L	
CARBON DIOXIDE	26		20-32 mmol/L	
CALCIUM	9.6		8.6-10.4 mg/dL	
PROTEIN, TOTAL	6.9		6.1-8.1 g/dL	
ALBUMIN	4.3		3.6-5.1 g/dL	
GLOBULIN	2.6		1.9-3.7 g/dL (calc)	
ALBUMIN/GLOBULIN RATIO	1.7		1.0-2.5 (calc)	
BILIRUBIN, TOTAL	0.7		0.2-1.2 mg/dL	
ALKALINE PHOSPHATASE	66		37-153 U/L	
AST	12		10-35 U/L	
ALT	7		6-29 U/L	

MAGNESIUM, RBC	4.6		4.0-6.4 mg/dL	AMD
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This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics Nichols Institute Chantilly, VA. It has not been cleared or approved by the U.S. Food and Drug Administration. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

PHOSPHATE (AS PHOSPHORUS)	3.8		2.5-4.5 mg/dL	TP
URIC ACID	5.3		2.5-7.0 mg/dL	TP
Therapeutic target for gout patients: <6.0 mg/dL				
LD	128		120-250 U/L	TP
GGT	12		3-70 U/L	TP
T4, FREE	1.2		0.8-1.8 ng/dL	TP
T3, FREE	3.7		2.3-4.2 pg/mL	TP
CBC (INCLUDES DIFF/PLT)				TP
WHITE BLOOD CELL COUNT	7.8		3.8-10.8 Thousand/uL	
RED BLOOD CELL COUNT	4.36		3.80-5.10 Million/uL	
HEMOGLOBIN	13.5		11.7-15.5 g/dL	
HEMATOCRIT	40.7		35.0-45.0 %	
MCV	93.3		80.0-100.0 fL	

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MCH	31.0		27.0-33.0 pg	
MCHC	33.2		32.0-36.0 g/dL	
RDW	11.8		11.0-15.0 %	
PLATELET COUNT	287		140-400 Thousand/uL	
MPV	10.8		7.5-12.5 fL	
ABSOLUTE NEUTROPHILS	4103		1500-7800 cells/uL	
ABSOLUTE LYMPHOCYTES	2972		850-3900 cells/uL	
ABSOLUTE MONOCYTES	593		200-950 cells/uL	
ABSOLUTE EOSINOPHILS	94		15-500 cells/uL	
ABSOLUTE BASOPHILS	39		0-200 cells/uL	
NEUTROPHILS	52.6		%	
LYMPHOCYTES	38.1		%	
MONOCYTES	7.6		%	
EOSINOPHILS	1.2		%	
BASOPHILS	0.5		%	
IRON, TOTAL	121		45-160 mcg/dL	TP
FERRITIN	54		16-232 ng/mL	TP

PERFORMING SITE:

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