

ODEN,JAMIE

DOB: 01/12/1979
Sex: F
Phone: (505) 306-1415
Patient ID: KBJ1982

Age: 45
Fasting: Y

Specimen: OZ185447D
Requisition: 0104500
Lab Reference ID: 00003938
Report Status: FINAL / SEE REPORT

Collected: 01/19/2024 07:57
Received: 01/19/2024 07:58
Reported: 01/24/2024 14:58

Client #: 31005802
JONES-SHORR,KAREN
JONES ND, KAREN (W)
960 LIBERTY ST SE
SALEM, OR 97302-4171
Phone: (503) 990-8395
Fax: (844) 778-7077

FASTING: YES

▲ LIPID PANEL, STANDARD

Analyte	Value	
▲ CHOLESTEROL, TOTAL	315 H	Reference Range: <200 mg/dL
HDL CHOLESTEROL	50	Reference Range: > OR = 50 mg/dL
TRIGLYCERIDES	119	Reference Range: <150 mg/dL

▲ LDL-CHOLESTEROL**239 H** mg/dL (calc)

LDL-C levels > or = 190 mg/dL may indicate familial hypercholesterolemia (FH). Clinical assessment and measurement of blood lipid levels should be considered for all first degree relatives of patients with an FH diagnosis. LDL Cholesterol (LDL-C) levels > or = 300 mg/dL may indicate homozygous familial hypercholesterolemia (HoFH). Untreated, these extremely high LDL-C levels can result in premature CV events and mortality. Patients should be identified early and provided appropriate interventions to reduce the cumulative LDL-C burden from birth.

For questions about testing for familial hypercholesterolemia, please call Quest Genomics Client Services at 1.866.GENE.INFO.

Jacobson T, et al. J National Lipid Association Recommendations for Patient-Centered Management of Dyslipidemia: Part 1 Journal of Clinical Lipidology 2015;9(2), 129-169.
Cuchel, M. et al. (2014). Homozygous familial hypercholesterolaemia: new insights and guidance for clinicians to improve detection and clinical management. European Heart Journal, 35(32), 2146-2157.
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**6.3 H** Reference Range: <5.0 (calc)**▲ NON HDL CHOLESTEROL****265 H** Reference Range: <130 mg/dL (calc)

Non-HDL level > or = 220 is very high and may indicate genetic familial hypercholesterolemia (FH). Clinical assessment and measurement of blood lipid levels should be considered for all first-degree relatives of patients with an FH diagnosis.

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.

▲ VITAMIN D,25-OH,TOTAL,IA

Analyte	Value
▲ VITAMIN D,25-OH,TOTAL,IA	28 L Reference Range: 30-100 ng/mL
Vitamin D Status	25-OH Vitamin D:
Deficiency:	<20 ng/mL
Insufficiency:	20 - 29 ng/mL
Optimal:	> or = 30 ng/mL
For 25-OH Vitamin D testing on patients on D2-supplementation and patients for whom quantitation of D2 and D3 fractions is required, the QuestAssured(TM) 25-OH VIT D, (D2,D3), LC/MS/MS is recommended: order code 92888 (patients >2yrs).	
COMMENT	
See Note 1	
Note 1	
For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ199 (This link is being provided for informational/ educational purposes only.)	

COMPREHENSIVE METABOLIC PANEL

Analyte	Value
GLUCOSE	93 Reference Range: 65-99 mg/dL
Fasting reference interval	
UREA NITROGEN (BUN)	13 Reference Range: 7-25 mg/dL
CREATININE	0.71 Reference Range: 0.50-0.99 mg/dL
EGFR	107 Reference Range: > OR = 60 mL/min/1.73m2
BUN/CREATININE RATIO	SEE NOTE: Reference Range: 6-22 (calc)
Not Reported: BUN and Creatinine are within reference range.	
SODIUM	139 Reference Range: 135-146 mmol/L
POTASSIUM	4.4 Reference Range: 3.5-5.3 mmol/L
CHLORIDE	108 Reference Range: 98-110 mmol/L
CARBON DIOXIDE	24 Reference Range: 20-32 mmol/L
CALCIUM	9.1 Reference Range: 8.6-10.2 mg/dL
PROTEIN, TOTAL	7.1 Reference Range: 6.1-8.1 g/dL
ALBUMIN	4.5 Reference Range: 3.6-5.1 g/dL
GLOBULIN	2.6 Reference Range: 1.9-3.7 g/dL (calc)
ALBUMIN/GLOBULIN RATIO	1.7 Reference Range: 1.0-2.5 (calc)
BILIRUBIN, TOTAL	0.8 Reference Range: 0.2-1.2 mg/dL
ALKALINE PHOSPHATASE	53 Reference Range: 31-125 U/L
AST	13 Reference Range: 10-35 U/L
ALT	13 Reference Range: 6-29 U/L

ANTI-MULLERIAN HORMONE (AMH), FEMALE

Analyte	Value
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ANTI-MULLERIAN HORMONE (AMH), FEMALE	0.93	Reference Range: 0.01-2.99 ng/mL
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CBC (INCLUDES DIFF/PLT)

Analyte	Value	
WHITE BLOOD CELL COUNT	5.6	Reference Range: 3.8-10.8 Thousand/uL
RED BLOOD CELL COUNT	4.98	Reference Range: 3.80-5.10 Million/uL
HEMOGLOBIN	14.7	Reference Range: 11.7-15.5 g/dL
HEMATOCRIT	44.5	Reference Range: 35.0-45.0 %
MCV	89.4	Reference Range: 80.0-100.0 fL
MCH	29.5	Reference Range: 27.0-33.0 pg
MCHC	33.0	Reference Range: 32.0-36.0 g/dL
RDW	12.8	Reference Range: 11.0-15.0 %
PLATELET COUNT	325	Reference Range: 140-400 Thousand/uL
MPV	10.3	Reference Range: 7.5-12.5 fL
ABSOLUTE NEUTROPHILS	3175	Reference Range: 1500-7800 cells/uL
ABSOLUTE LYMPHOCYTES	1674	Reference Range: 850-3900 cells/uL
ABSOLUTE MONOCYTES	521	Reference Range: 200-950 cells/uL
ABSOLUTE EOSINOPHILS	190	Reference Range: 15-500 cells/uL
ABSOLUTE BASOPHILS	39	Reference Range: 0-200 cells/uL
NEUTROPHILS	56.7	%
LYMPHOCYTES	29.9	%
MONOCYTES	9.3	%
EOSINOPHILS	3.4	%
BASOPHILS	0.7	%

FERRITIN

Analyte	Value	
FERRITIN	38	Reference Range: 16-232 ng/mL

INSULIN

Analyte	Value	
INSULIN	17.8	uIU/mL

Reference Range < or = 18.4

Risk:
 Optimal < or = 18.4
 Moderate NA
 High >18.4

Adult cardiovascular event risk category cut points (optimal, moderate, high) are based on Insulin Reference Interval studies performed at Quest Diagnostics in 2022.

PROGESTERONE

Analyte	Value
PROGESTERONE	10.9 ng/mL
Reference Ranges	
Female	
Follicular Phase	< 1.0
Luteal Phase	2.6-21.5
Post menopausal	< 0.5
Pregnancy	
1st Trimester	4.1-34.0
2nd Trimester	24.0-76.0
3rd Trimester	52.0-302.0

T4, FREE

Analyte	Value
T4, FREE	1.0 Reference Range: 0.8-1.8 ng/dL

TSH

Analyte	Value
TSH	2.99 mIU/L
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

ESTRADIOL

Analyte	Value
ESTRADIOL	62 pg/mL
Reference Range	
Follicular Phase:	19-144
Mid-Cycle:	64-357
Luteal Phase:	56-214
Postmenopausal:	< or = 31

Reference range established on post-pubertal patient population. No pre-pubertal reference range established using this assay. For any patients for whom low Estradiol levels are anticipated (e.g. males, pre-pubertal children and hypogonadal/post-menopausal females), the Quest Diagnostics Nichols Institute Estradiol, Ultrasensitive, LCMSMS assay is recommended (order code 30289).

Please note: patients being treated with the drug fulvestrant (Faslodex(R)) have demonstrated significant interference in immunoassay methods for estradiol measurement. The cross reactivity could lead to falsely elevated estradiol test results leading to an inappropriate clinical assessment of estrogen status. Quest Diagnostics order code 30289-Estradiol, Ultrasensitive LC/MS/MS demonstrates negligible cross reactivity with fulvestrant.

T3, FREE


Analyte	Value
T3, FREE	2.9 Reference Range: 2.3-4.2 pg/mL

Performing Sites

EZ Quest Diagnostics/Nichols SJC-San Juan Capistrano,, 33608 Ortega Hwy, San Juan Capistrano, CA 92675-2042 Laboratory Director: Irina Maramica MD,PhD,MBA

NW Quest Diagnostics-Seattle 1737 Airport, 1737 Airport Way S, Suite 200, Seattle, WA 98134-1636 Laboratory Director: Roger Graham

Key

 Priority Out of Range  Out of Range

These results have been sent to the person who ordered the tests. Your receipt of these results should not be viewed as medical advice and is not meant to replace discussion with your doctor or other healthcare professional.

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