

CHAMBERS,KEVIN

DOB: 09/14/1993
Sex: M
Phone: (505) 967-8903
Patient ID: 4094481

Age: 31
Fasting:

Specimen: JZ492660A
Requisition: 0001967
Lab Reference ID: 45850782
Report Status: FINAL / SEE REPORT

Collected: 03/18/2025 08:10
Received: 03/18/2025 08:12
Reported: 03/25/2025 08:11

Client #: 50544437
JAY,JAN
ENHANCED WELLNESS
JAN JAY DOM/JOSEPH JAROS MD
5200 EUBANK BLVD NE STE C3
ALBUQUERQUE, NM 87111-1764
Phone: (505) 323-8100
Fax: (505) 292-0555

▲ **LIPID PANEL, STANDARD**

Analyte	Value	
CHOLESTEROL, TOTAL	195	Reference Range: <200 mg/dL
▲ HDL CHOLESTEROL	35 L	Reference Range: > OR = 40 mg/dL
▲ TRIGLYCERIDES	258 H	Reference Range: <150 mg/dL
If a non-fasting specimen was collected, consider repeat triglyceride testing on a fasting specimen if clinically indicated. Jacobson et al. J. of Clin. Lipidol. 2015;9:129-169.		
▲ LDL-CHOLESTEROL	120 H	mg/dL (calc)
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/HDLC RATIO	5.6 H	Reference Range: <5.0 (calc)
▲ NON HDL CHOLESTEROL	160 H	Reference Range: <130 mg/dL (calc)
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**

Analyte	Value	
▲ GLUCOSE	101 H	Reference Range: 65-99 mg/dL
Fasting reference interval		
For someone without known diabetes, a glucose value between 100 and 125 mg/dL is consistent with prediabetes and should be confirmed with a follow-up test.		
UREA NITROGEN (BUN)	19	Reference Range: 7-25 mg/dL
CREATININE	0.96	Reference Range: 0.60-1.26 mg/dL
EGFR	108	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	137	Reference Range: 135-146 mmol/L
POTASSIUM	4.2	Reference Range: 3.5-5.3 mmol/L

CHLORIDE	102	Reference Range: 98-110 mmol/L
CARBON DIOXIDE	29	Reference Range: 18-30 mmol/L
CALCIUM	9.5	Reference Range: 8.6-10.3 mg/dL
PROTEIN, TOTAL	7.5	Reference Range: 6.1-8.1 g/dL
ALBUMIN	4.9	Reference Range: 3.6-5.1 g/dL
GLOBULIN	2.6	Reference Range: 1.9-3.7 g/dL (calc)
ALBUMIN/GLOBULIN RATIO	1.9	Reference Range: 1.0-2.5 (calc)
BILIRUBIN, TOTAL	0.8	Reference Range: 0.2-1.2 mg/dL
ALKALINE PHOSPHATASE	83	Reference Range: 36-130 U/L
AST	25	Reference Range: 10-40 U/L
ALT	44	Reference Range: 9-46 U/L

▲ IRON AND TOTAL IRON BINDING CAPACITY

Analyte	Value	
IRON, TOTAL	153	Reference Range: 50-180 mcg/dL
▲ IRON BINDING CAPACITY	433 H	Reference Range: 250-425 mcg/dL (calc)
% SATURATION	35	Reference Range: 20-48 % (calc)

▲ CBC (INCLUDES DIFF/PLT)

Analyte	Value	
WHITE BLOOD CELL COUNT	5.5	Reference Range: 3.8-10.8 Thousand/uL
▲ RED BLOOD CELL COUNT	6.33 H	Reference Range: 4.33-5.82 Million/uL
▲ HEMOGLOBIN	19.4 H	Reference Range: 13.7-17.7 g/dL
Verified by repeat analysis.		
▲ HEMATOCRIT	56.4 H	Reference Range: 41.5-53.8 %
MCV	89.1	Reference Range: 80.0-100.0 fL
MCH	30.6	Reference Range: 27.0-33.0 pg
MCHC	34.4	Reference Range: 32.0-36.0 g/dL
For adults, a slight decrease in the calculated MCHC value (in the range of 30 to 32 g/dL) is most likely not clinically significant; however, it should be interpreted with caution in correlation with other red cell parameters and the patient's clinical condition.		
RDW	12.4	Reference Range: 11.0-15.0 %
PLATELET COUNT	221	Reference Range: 140-400 Thousand/uL
MPV	10.6	Reference Range: 7.5-12.5 fL
ABSOLUTE NEUTROPHILS	3003	Reference Range: 1500-7800 cells/uL
ABSOLUTE LYMPHOCYTES	1771	Reference Range: 850-3900 cells/uL
ABSOLUTE MONOCYTES	539	Reference Range: 200-950 cells/uL
ABSOLUTE EOSINOPHILS	149	Reference Range: 15-500 cells/uL
ABSOLUTE BASOPHILS	39	Reference Range: 0-200 cells/uL

NEUTROPHILS	54.6	%
LYMPHOCYTES	32.2	%
MONOCYTES	9.8	%
EOSINOPHILS	2.7	%
BASOPHILS	0.7	%

VITAMIN D,25-OH,TOTAL,IA

Analyte	Value
VITAMIN D,25-OH,TOTAL,IA	30 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.)

TESTOSTERONE, FREE (DIALYSIS), TOTAL (MS) AND SEX HORMONE BINDING GLOBULIN

Analyte	Value
TESTOSTERONE, TOTAL, MS	379 Reference Range: 250-1100 ng/dL
Men with clinically significant hypogonadal symptoms and testosterone values repeatedly in the range of the 200-300 ng/dL or less, may benefit from testosterone treatment after adequate risk and benefits counseling.	
For additional information, please refer to https://education.questdiagnostics.com/faq/FAQ165 (This link is being provided for informational/educational purposes only.) (Note)	
This test was developed and its analytical performance characteristics have been determined by medfusion. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.	
TESTOSTERONE, FREE	81.5 Reference Range: 35.0-155.0 pg/mL
(Note)	
This test was developed and its analytical performance characteristics have been determined by medfusion. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.	
MDF med fusion 2501 South State Highway 121,Suite 1100 Lewisville TX 75067 972-966-7300 Ithiel James L. Frame, MD, PhD	
SEX HORMONE BINDING GLOBULIN	18.2 Reference Range: 10-50 nmol/L

MAGNESIUM

Analyte	Value
MAGNESIUM	2.2 Reference Range: 1.5-2.5 mg/dL

T3 REVERSE, LC/MS/MS

Analyte	Value
T3 REVERSE, LC/MS/MS	12 Reference Range: 8-25 ng/dL
<p>This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.</p>	

ANA SCREEN, IFA, W/REFL TITER AND PATTERN

Analyte	Value
ANA SCREEN, IFA	NEGATIVE Reference Range: NEGATIVE
<p>ANA IFA is a first line screen for detecting the presence of up to approximately 150 autoantibodies in various autoimmune diseases. A negative ANA IFA result suggests an ANA-associated autoimmune disease is not present at this time, but is not definitive. If there is high clinical suspicion for Sjogren's syndrome, testing for anti-SS-A/Ro antibody should be considered. Anti-Jo-1 antibody should be considered for clinically suspected inflammatory myopathies.</p> <p>AC-0: Negative</p> <p>International Consensus on ANA Patterns (https://doi.org/10.1515/cclm-2018-0052)</p> <p>For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ177 (This link is being provided for informational/ educational purposes only.)</p>	

HS CRP

Analyte	Value
HS CRP	2.7 mg/L
<p>Reference Range Optimal <1.0 Jellinger PS et al. Endocr Pract.2017;23(Suppl 2):1-87. For ages >17 Years: hs-CRP mg/L Risk According to AHA/CDC Guidelines <1.0 Lower relative cardiovascular risk. 1.0-3.0 Average relative cardiovascular risk. 3.1-10.0 Higher relative cardiovascular risk. Consider retesting in 1 to 2 weeks to exclude a benign transient elevation in the baseline CRP value secondary to infection or inflammation. >10.0 Persistent elevation, upon retesting, may be associated with infection and inflammation.</p> <p>Pearson TA, Mensah GA, Alexander RW, et al. Markers of inflammation and cardiovascular disease: application to clinical and public health practice: A statement for healthcare professionals from the Centers for Disease Control and Prevention and the American Heart Association. Circulation 2003; 107(3): 499-511.</p>	

GLUCOSE-6-PHOSPHATE DEHYDROGENASE, QN

Analyte	Value
GLUCOSE-6-PHOSPHATE DEHYDROGENASE	12.1 Reference Range: 7.0-20.5 U/g Hgb

CORTISOL, TOTAL, LC/MS

Analyte	Value
CORTISOL, TOTAL, LC/MS	21.5 mcg/dL
Adult Reference Ranges for Cortisol, Total:	
8-10 AM 4.6-20.6 mcg/dL	
4-6 PM 1.8-13.6 mcg/dL	
Cortisol Response to ACTH	
Peak >20.0 mcg/dL	
Peak >16.0 mcg/dL after IM injection	
This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.	

THYROID PEROXIDASE ANTIBODIES

Analyte	Value
THYROID PEROXIDASE ANTIBODIES	<1 Reference Range: <9 IU/mL

HOMOCYSTEINE

Analyte	Value
HOMOCYSTEINE	6.5 Reference Range: <11.4 umol/L
Homocysteine is increased by functional deficiency of folate or vitamin B12. Testing for methylmalonic acid differentiates between these deficiencies. Other causes of increased homocysteine include renal failure, folate antagonists such as methotrexate and phenytoin, and exposure to nitrous oxide.	
Selhub J, et al., Ann Intern Med. 1999;131(5):331-9.	

DHEA SULFATE

Analyte	Value
DHEA SULFATE	170 Reference Range: 93-415 mcg/dL

FERRITIN

Analyte	Value
FERRITIN	114 Reference Range: 38-380 ng/mL

PROGESTERONE

Analyte	Value
PROGESTERONE	<0.5 Reference Range: <1.4 ng/mL

T3, TOTAL

Analyte	Value
T3, TOTAL	105 Reference Range: 76-181 ng/dL

T4, FREE

Analyte	Value
T4, FREE	1.1 Reference Range: 0.8-1.8 ng/dL

T4 (THYROXINE), TOTAL

Analyte	Value
T4 (THYROXINE), TOTAL	6.2 Reference Range: 4.9-10.5 mcg/dL

TSH

Analyte	Value
TSH	2.19 Reference Range: 0.40-4.50 mIU/L

T3, FREE

Analyte	Value
T3, FREE	3.9 Reference Range: 2.3-4.2 pg/mL

ESTRADIOL

Analyte	Value
ESTRADIOL 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.	29 Reference Range: < OR = 39 pg/mL

VITAMIN B12/FOLATE, SERUM PANEL

Analyte	Value
VITAMIN B12	746 Reference Range: 200-1100 pg/mL
FOLATE, SERUM Reference Range Low: <3.4 Borderline: 3.4-5.4 Normal: >5.4	19.4 ng/mL

PSA (FREE AND TOTAL)

Analyte	Value
PSA, TOTAL	0.6 Reference Range: < OR = 4.0 ng/mL
PSA, FREE	0.3 ng/mL

PSA, % FREE**50** Reference Range: >25 % (calc)

PSA(ng/mL)	Free PSA(%)	Estimated(x) Probability of Cancer(as%)
0-2.5	(*)	Approx. 1
2.6-4.0(1)	0-27(2)	24(3)
4.1-10(4)	0-10	56
	11-15	28
	16-20	20
	21-25	16
	>or =26	8
>10(+)	N/A	>50

References:(1)Catalona et al.:Urology 60: 469-474 (2002)
(2)Catalona et al.:J.Urol 168: 922-925 (2002)
Free PSA(%) Sensitivity(%) Specificity(%)
< or = 25 85 19
< or = 30 93 9
(3)Catalona et al.:JAMA 277: 1452-1455 (1997)
(4)Catalona et al.:JAMA 279: 1542-1547 (1998)

(x)These estimates vary with age, ethnicity, family history and DRE results.
(*)The diagnostic usefulness of % Free PSA has not been established in patients with total PSA below 2.6 ng/mL
(+)In men with PSA above 10 ng/mL, prostate cancer risk is determined by total PSA alone.

The Total PSA value from this assay system is standardized against the equimolar PSA standard. The test result will be approximately 20% higher when compared to the WHO-standardized Total PSA (Siemens assay). Comparison of serial PSA results should be interpreted with this fact in mind.

PSA was performed using the Beckman Coulter Immunoassay method. Values obtained from different assay methods cannot be used interchangeably. PSA levels, regardless of value, should not be interpreted as absolute evidence of the presence or absence of disease.

HEMOGLOBIN A1c**Analyte** **Value****HEMOGLOBIN A1c****5.4** Reference Range: <5.7 % of total Hgb

For the purpose of screening for the presence of diabetes:

<5.7% Consistent with the absence of diabetes
5.7-6.4% Consistent with increased risk for diabetes (prediabetes)
> or =6.5% Consistent with diabetes

This assay result is consistent with a decreased risk of diabetes.

Currently, no consensus exists regarding use of hemoglobin A1c for diagnosis of diabetes in children.

According to American Diabetes Association (ADA) guidelines, hemoglobin A1c <7.0% represents optimal control in non-pregnant diabetic patients. Different metrics may apply to specific patient populations. Standards of Medical Care in Diabetes(ADA).

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

IG Quest Diagnostics-Dallas Lab, 4770 Regent Blvd, Irving, TX 75063-2445 Laboratory Director: Clare McCormick-Baw MD, PhD

QJP Quest Diagnostics-Central Laboratory, 5601 Office Blvd NE, Albuquerque, NM 87109-5879 Laboratory Director: Robert L Breckenridge, MD

Z3E MedFusion-MedFusion, 2501 South State Highway 121, Suite 1100, Lewisville, TX 75067-8188 Laboratory Director: Ithiel James L Frame MD,PhD

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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