



Specimen: Requisition: 0000430 Lab Reference ID: 011230 Report Status: FINAL / SEE REPORT Collected: 03/05/2025 06:07 Received: 03/05/2025 06:08 Reported: 03/14/2025 14:06 Client #:

DAMASCO,LEO

JASONHEALTH.COM

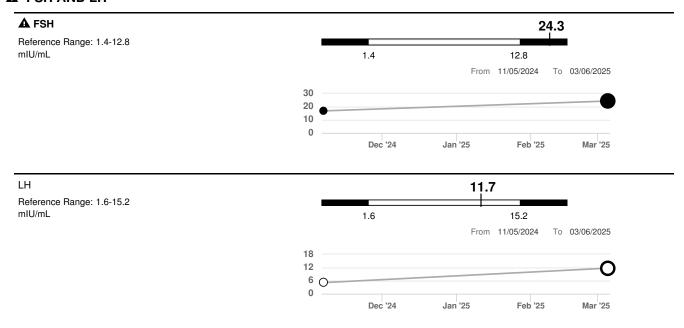
1887 WHITNEY MESA DR # 3040

HENDERSON, NV 89014-2069

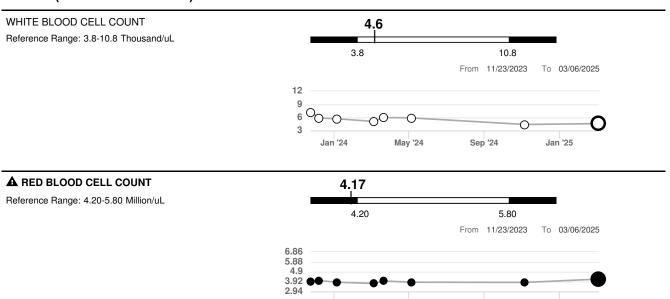
Phone: (702) 935-0035

### FASTING:YES

### **▲ FSH AND LH**



# **▲** CBC (INCLUDES DIFF/PLT)

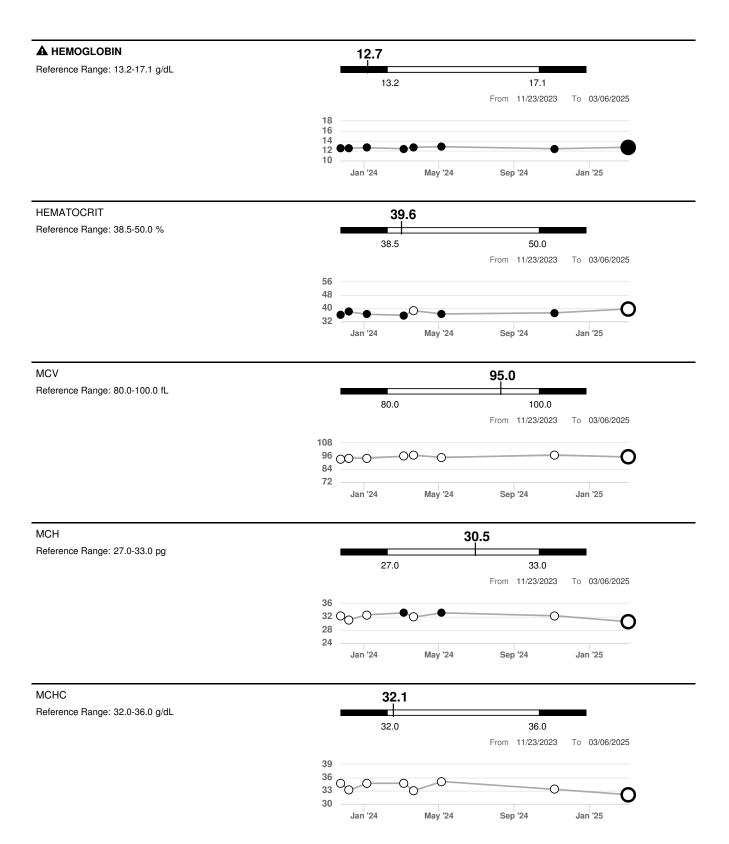


Jan '24

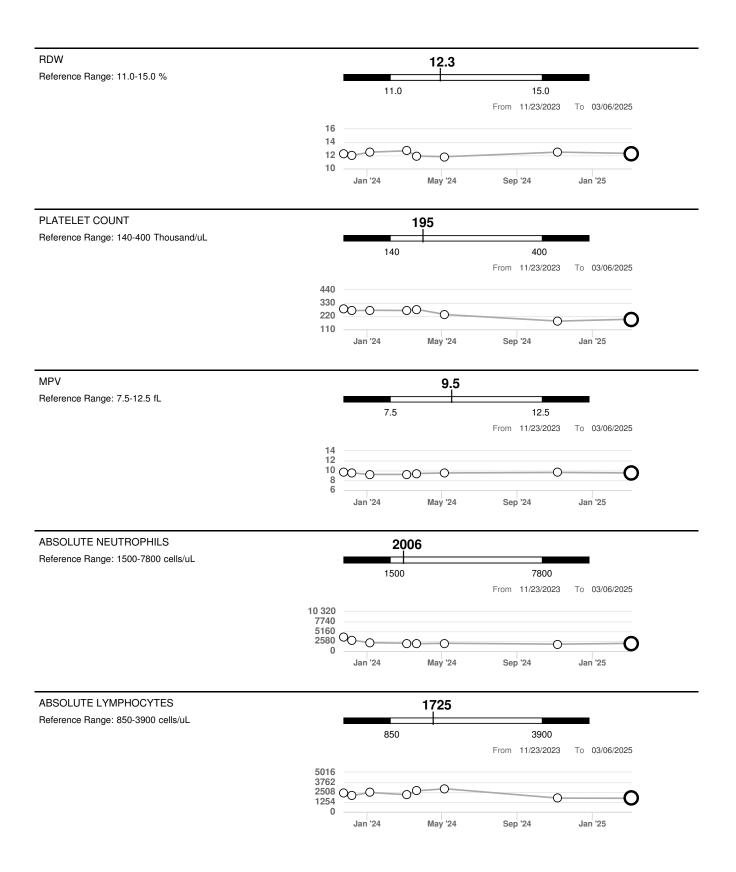
May '24

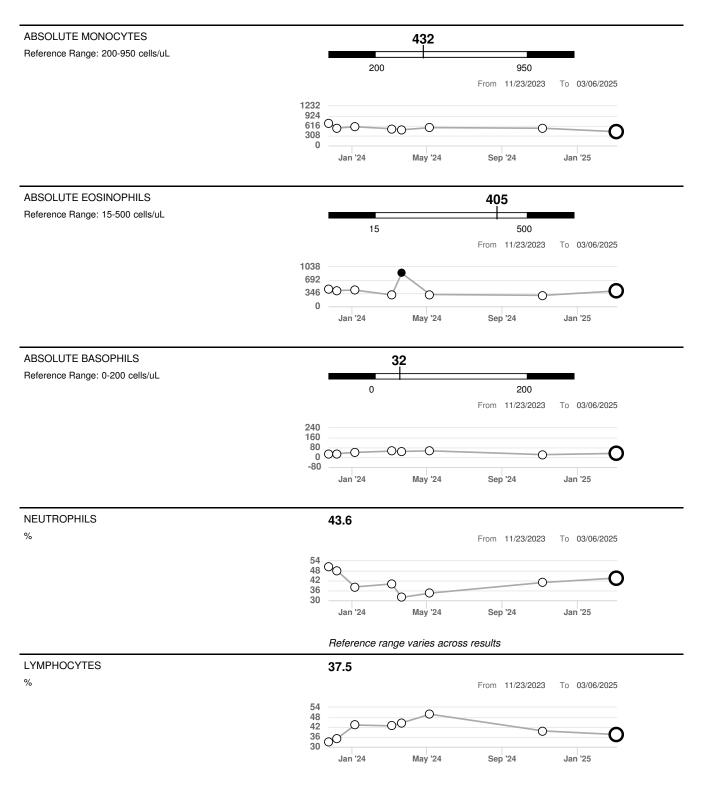
Sep '24

Jan '25

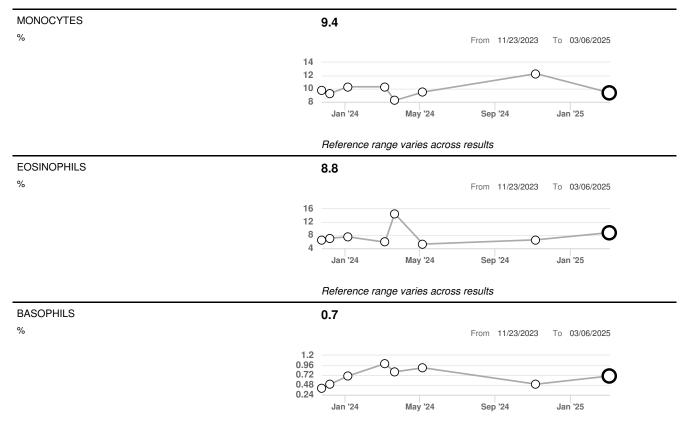


For adults, a slight decrease in the calculated MCHC value (in the range of 30 to 32 g/dL) is most likely not clinically significant; however, it should be interpreted with caution in correlation with other red cell parameters and the patient's clinical condition.





Reference range varies across results



Reference range varies across results

### ▲ CANDIDA ALBICANS AB (IGG,IGA,IGM)

▲ C.ALBICANS IGG	1.2 H
	No Historical Data
▲ C.ALBICANS IGA	2.2 H
	No Historical Data
C.ALBICANS IGM	0.6

No Historical Data

REFERENCE RANGE: <1.0

INTERPRETIVE CRITERIA:

Antibody Not Detected <1.0 > or = 1.0Antibody Detected

Systemic candidiasis is often characterized by markedly elevated levels of IgG, IgA, and IgM recognizing Candida. However, interpretation of Candida antibody results is complicated by antibody detection in approximately 40% of healthy individuals and up to 70% of patients positive for other fungal antibodies. Further, antibody responses may be blunted in immunocompromised patients at risk for systemic candidiasis. Candida antibody levels should be considered within the context of clinical findings and results from other relevant laboratory tests, such as Candida antigen detection and/or culture.

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

### **COMPREHENSIVE METABOLIC PANEL**

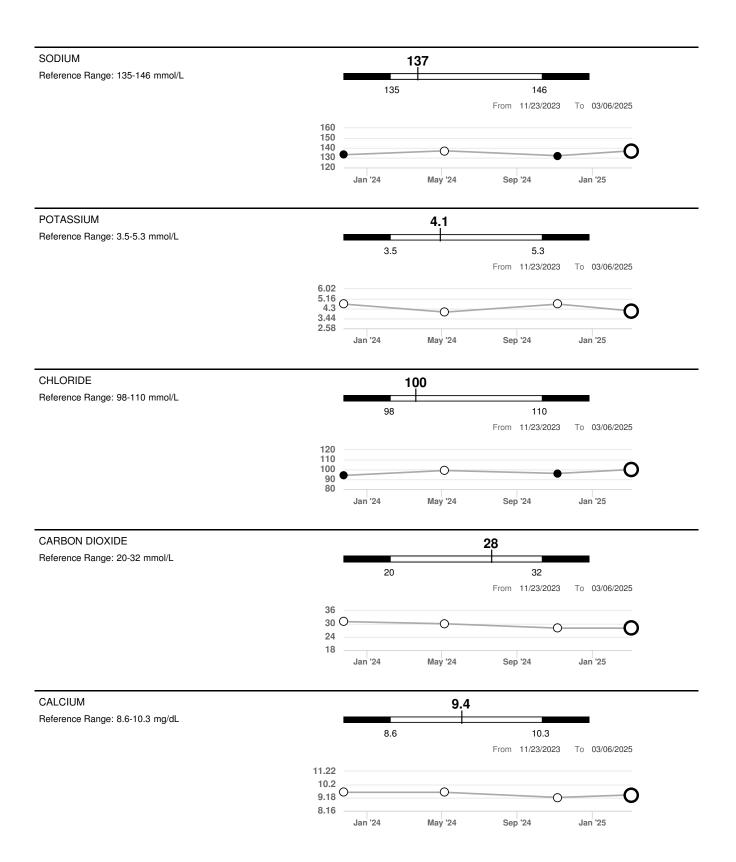
#### GLUCOSE 85 Reference Range: 65-99 mg/dL 65 99 From 11/23/2023 To 03/06/2025 144 120 96 72 48 Jan '24 May '24 Sep '24 Jan '25 Fasting reference interval UREA NITROGEN (BUN) 19 Reference Range: 7-25 mg/dL 25 From 11/23/2023 To 03/06/2025 32 24 16 8 0 Jan '24 May '24 Sep '24 Jan '25 **CREATININE** 0.82 Reference Range: 0.70-1.28 mg/dL 0.70 1.28 From 11/23/2023 To 03/06/2025 1.56 1.3 1.04 0.78 0.52 0 Jan '24 May '24 Sep '24 Jan '25 **EGFR** 94 Reference Range: > OR = 60 mL/min/1.73m2 From 11/23/2023 To 03/06/2025 112 96 80 O 64 48 Jan '24 May '24 Sep '24 Jan '25

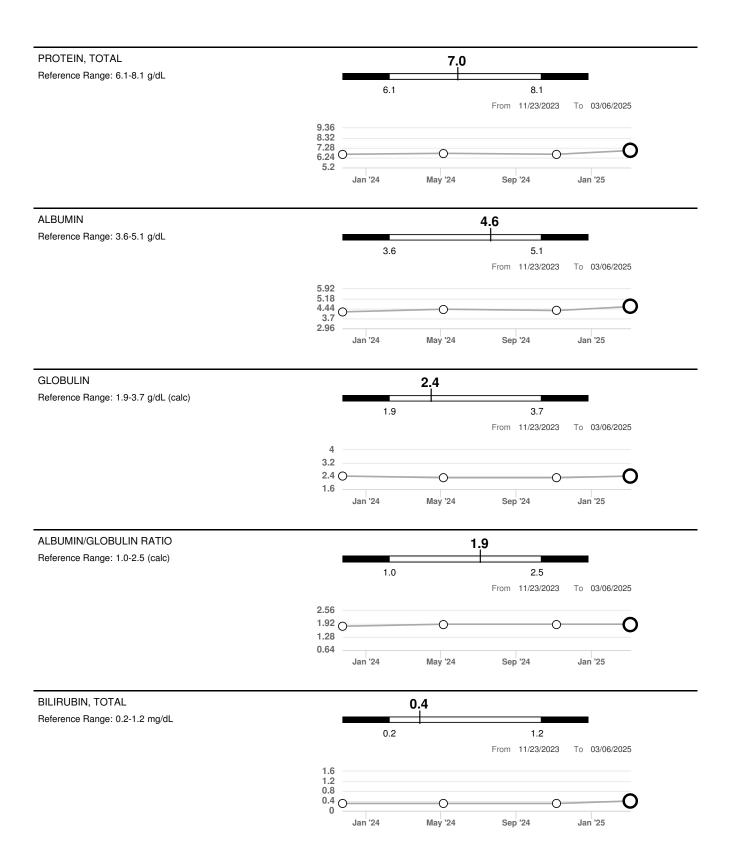
BUN/CREATININE RATIO

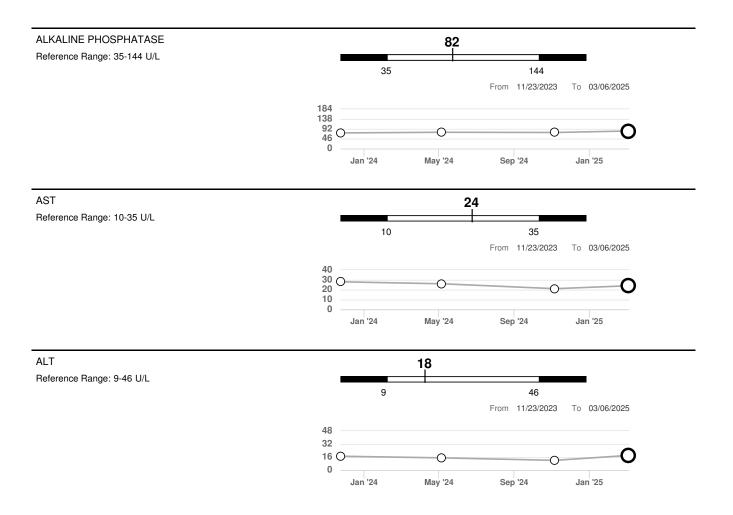
Reference Range: 6-22 (calc)

Not Reported:  $\ensuremath{\mathsf{BUN}}$  and  $\ensuremath{\mathsf{Creatinine}}$  are within reference range.

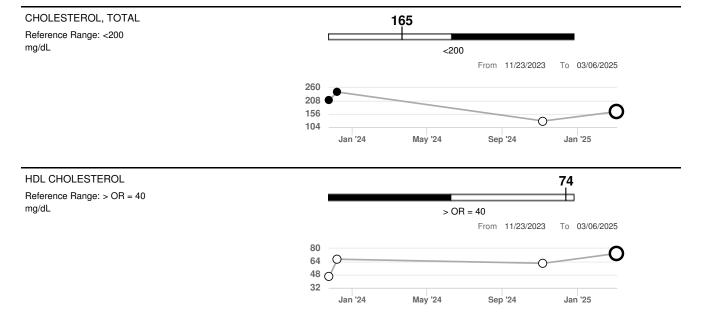
SEE NOTE:

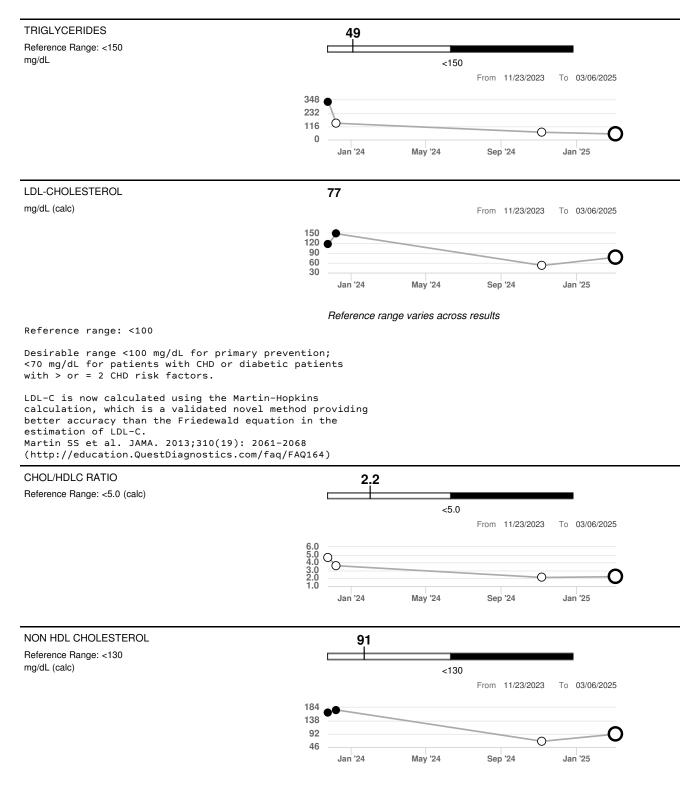






# LIPID PANEL, STANDARD





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.

### **HS CRP**

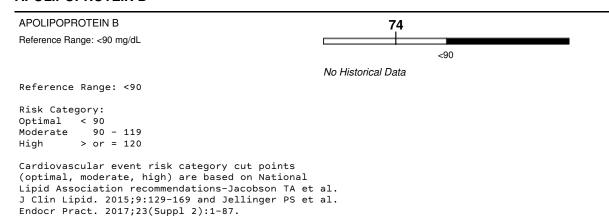


Reference Range Optimal <1.0 Jellinger PS et al. Endocr Pract.2017;23(Suppl 2):1-87. For ages >17 Years: hs-CRP mg/L Risk According to AHA/CDC Guidelines Lower relative cardiovascular risk. <1.0 1.0-3.0 Average relative cardiovascular risk. 3.1-10.0 Higher relative cardiovascular risk. Consider retesting 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. 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

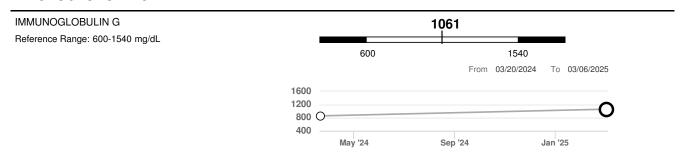
Centers for Disease Control and Prevention and the American Heart Association. Circulation 2003; 107(3):

### APOLIPOPROTEIN B

499-511.



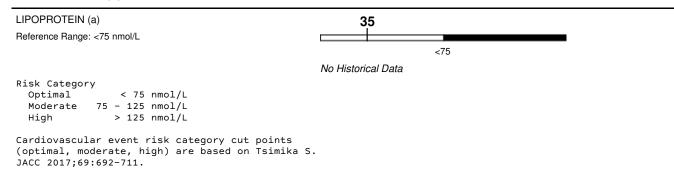
## **IMMUNOGLOBULIN G**



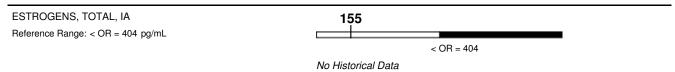
### **CREATINE KINASE, TOTAL**



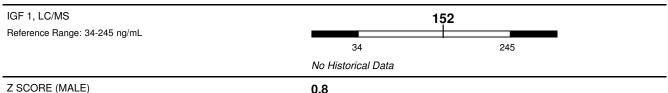
### LIPOPROTEIN (a)



### **ESTROGENS, TOTAL, IA**



### IGF 1, LC/MS



8.0

Reference Range: -2.0 - +2.0 SD No Historical Data

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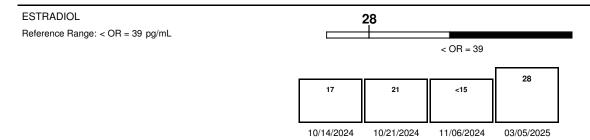
### LP PLA2 ACTIVITY



Risk: Optimal <=123 nmol/min/mL; High >123 nmol/min/mL.

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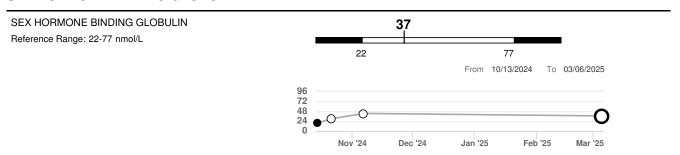
### **ESTRADIOL**



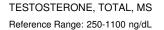
Reference range established on post-pubertal patient population. No pre-pubertal reference range established using this assay. For any patients for whom low Estradiol levels are anticipated (e.g. males, pre-pubertal children and hypogonadal/post-menopausal females), the Quest Diagnostics Nichols Institute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

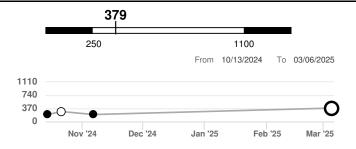
Please note: patients being treated with the drug fulvestrant (Faslodex(R)) have demonstrated significant interference in immunoassay methods for estradiol measurement. The cross reactivity could lead to falsely elevated estradiol test results leading to an inappropriate clinical assessment of estrogen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

#### **SEX HORMONE BINDING GLOBULIN**



### TESTOSTERONE, FREE (DIALYSIS) AND TOTAL, MS





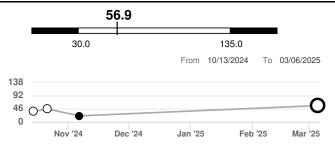
Men with clinically significant hypogonadal symptoms and testosterone values repeatedly in the range of the 200-300 ng/dL or less, may benefit from testosterone treatment after adequate risk and benefits counseling.

For additional information, please refer to http://education.questdiagnostics.com/faq/ TotalTestosteroneLCMSMSFAQ165 (This link is being provided for informational/ educational purposes only.)

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.

#### TESTOSTERONE, FREE

Reference Range: 30.0-135.0 pg/mL



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.

### **Performing Sites**

AMD Quest Diagnostics/Nichols Chantilly-Chantilly VA, 14225 Newbrook Dr, Chantilly, VA 20151-2228 Laboratory Director: Patrick W Mason M.D.,PhD

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

TP Quest Diagnostics-Tampa, 4225 E Fowler Ave, Tampa, FL 33617-2026 Laboratory Director: Weston H Rothrock MD

#### Key

Priority Out of Range Out of Range

Note: Data displayed only for results that meet strict identification matching. Historical result view may vary based on corrected or updated patient demographics. The reference range displayed may vary due to potential changes in laboratory testing methods. Please refer to the published reference range on each lab report.

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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