

**Specimen Collected: 09-Mar-21 05:42**

**Vedolizumab Quantitation with Antibodies** | **Received: 09-Mar-21 10:11** | **Report/Verified: 09-Mar-21 09:14**

Procedure	Result	Units	Reference Interval
Vedolizumab Quantitation, Serum	<2.0 <sup>L</sup> <sup>f1</sup>	mcg/mL	
Vedolizumab Antibodies, Serum	9.8 <sup>H</sup>	ng/mL	<9.8
Vedolizumab Interpretation	SEE NOTE <sup>f2</sup>		

**Result Footnote**

f1: Vedolizumab Quantitation, Serum  
 -----REFERENCE VALUE-----  
 Lower limit of quantitation = 2.0 mcg/mL  
 -----ADDITIONAL INFORMATION-----  
 This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

f2: Vedolizumab Interpretation  
 RESULT: Presence of antibody-to-vedolizumab detected.  
 -----ADDITIONAL INFORMATION-----  
 This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.  
 Test Performed by:  
 Mayo Clinic Laboratories - Rochester Superior Drive  
 3050 Superior Drive NW, Rochester, MN 55901  
 Lab Director: William G. Morice M.D. Ph.D.; CLIA# 24D1040592

\*=Abnormal, #=Corrected, C=Critical, f=Result Footnote, H-High, i-Test Information, L-Low, t-Interpretive Text, @=Performing lab

*Unless otherwise indicated, testing performed at:*

**ARUP Laboratories**

500 Chipeta Way, Salt Lake City, UT 84108

Laboratory Director: Tracy I. George, MD

**ARUP Accession:** 21-068-900005

**Report Request ID:** 14707729

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