

Specimen Collected: 22-Jun-21 17:33

Rheumatoid Arthritis Pan		Received: 22-Jun-21 17:44	Report/Verified: 22-Jun-21 18:12
Procedure	Result	Units	Reference Interval
Carbamylated Protein Antibody, IgG	20 ^H ⁱ¹	Units	0-19
Rheumatoid Arthritis Pan		Received: 22-Jun-21 17:44	Report/Verified: 23-Jun-21 11:02
Procedure	Result	Units	Reference Interval
Cyclic Citrullinated Peptide Ab, IgG	20 ^H ⁱ²	Units	0-19
Rheumatoid Arthritis Pan		Received: 22-Jun-21 17:44	Report/Verified: 23-Jun-21 12:48
Procedure	Result	Units	Reference Interval
Rheumatoid Factor	15 ^H	IU/mL	0-14

Test Information

i1: Carbamylated Protein Antibody, IgG

INTERPRETIVE INFORMATION: Carbamylated Protein (CarP) Ab, IgG

Anti-carbamylated protein (anti-CarP) IgG antibodies are present in about 34-53 percent of patients with RA, have specificities of greater than 90 percent and can occur in RA patients seronegative for both rheumatoid factor and anti-CCP. These autoantibodies may be present in the preclinical phase of disease, are associated with future RA development, and may predict radiographic joint destruction. Patients with weak positive results should be monitored and testing repeated.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

i2: Cyclic Citrullinated Peptide Ab, IgG

INTERPRETIVE INFORMATION: Cyclic Citrullinated Peptide Antibody, IgG

19 Units or less Negative
 20-39 Units Weak Positive
 40-59 Units Moderate Positive
 60 Units or greater Strong Positive

Anti-cyclic citrullinated peptide (anti-CCP), IgG antibodies are present in about 69-83 percent of patients with rheumatoid arthritis (RA) and have specificities of 93-95 percent. These autoantibodies may be present in the preclinical phase of disease, are associated with future RA development, and may predict radiographic joint destruction. Patients with weak positive results should be monitored and testing repeated.

*=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: 21-173-900278

Report Request ID: 15025139

Printed: 23-Jun-21 15:04