

Overview

Useful For

Assisting in the diagnostic process of ankylosing spondylitis, juvenile rheumatoid arthritis, and Reiter syndrome

Method Name

FlowCytometry

NY State Available

Yes

Specimen

Specimen Type

Whole Blood EDTA

Specimen Required

Specimen must arrive within 96 hours of draw.

Container/Tube: Lavender top (EDTA)

Specimen Volume: 6 mL

Collection Instructions: Do not transfer blood to other containers.

Forms

If not ordering electronically, complete, print, and send a [General Request](#) (T239) with the specimen.

Specimen Minimum Volume

1 mL

Reject Due To

All specimens will be evaluated at Mayo Clinic Laboratories for test suitability.

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Whole Blood EDTA	Ambient	4 days	

Clinical and Interpretive

Clinical Information

This major histocompatibility coded class I antigen is associated with ankylosing spondylitis, juvenile rheumatoid arthritis, and Reiter syndrome. The mechanism of the association is not understood but probably is that of linkage disequilibrium.

There is an increased prevalence of HLA-B27 in certain rheumatic diseases, particularly ankylosing spondylitis.

Studies have demonstrated that the B*27:06 allele, which is present in a small percentage of individuals of Asian ethnicity, may not be associated with ankylosing spondylitis.

Reference Values

An interpretive report will be provided.

Interpretation

Approximately 8% of the normal population carries the HLA-B27 antigen.

HLA-B27 is present in approximately 89% of patients with ankylosing spondylitis, 79% of patients with Reiter syndrome, and 42% of patients with juvenile rheumatoid arthritis. However, lacking other data, it is not diagnostic for these disorders.

Cautions

Orders received for both this test and SSO1 / HLA Class I Molecular Phenotype, Blood or DISI / [HLA Class I Molecular Typing Disease Association](#) (which provides data on all HLA Class I low-resolution antigens, including B27) will be questioned due to test overlap. This HLA-B27 test is best used alone if a particular disease such as ankylosing spondylitis is under consideration.

Extreme temperature changes during shipping may alter the specimen making it unacceptable for testing.

Clinical Reference

1. Brewerton DA, Hart FD, Nicholls A, et al: Ankylosing spondylitis and HLA-27. Lancet 1973;1:904-907
2. Albrecht J, Muller HA: HLA-B27 typing by use of flow cytofluorometry. Clin Chem 1987;33:1619-1623

Performance

Method Description

Anti-HLA fluorescein isothiocyanate (FITC)/CD3 phycoerythrin monoclonal antibody reagent is added to human whole blood. The fluorochrome-labeled antibodies bind specifically to leukocyte surface antigens. The stained specimens are treated with lysing solution to lyse red blood cells, then washed and fixed prior to flow cytometric analysis. The flow cytometer is set up using BD caliBRITE beads with Autocomp software and HLA-B27 calibration beads with the HLA-B27 software. The HLA-B27 software first identifies, on a forward scatter (FSC) versus fluorescence 2 (FL2) dot plot, the cluster of events with a uniformly bright CD3-positive signal (T-lymphocytes). During analysis, the median fluorescence intensity of the anti-HLA-B27 FITC signal is calculated for the events included in the FSC/FL2 gate. Specimens with a median fluorescence 1 channel result greater than or equal to the decision marker are considered HLA-B27 positive. Specimens with a median channel result lower than the decision marker are considered HLA-B27 negative. This decision marker is encoded in the suffix of the reagent lot number listed on the vial label. (Package insert: HLA-B27 Test Kit. Becton-Dickinson, San Jose, CA)

PDF Report

No

Day(s) and Time(s) Test Performed

Monday through Friday; 7:30-5 p.m.

Saturday; 10 a.m.-6 p.m.

Analytic Time

2 days

Maximum Laboratory Time

6 days

Specimen Retention Time

7 days

Performing Laboratory Location

Rochester

Fees and Codes**Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their Regional Manager. For assistance, contact [Customer Service](#).

Test Classification

This test has been cleared or approved by the U.S. Food and Drug Administration and is used per manufacturer's instructions. Performance characteristics were verified by Mayo Clinic in a manner consistent with CLIA requirements.

CPT Code Information

86812

LOINC® Information

Test ID	Test Order Name	Order LOINC Value
LY27B	HLA-B27, B	26028-1

Result ID	Test Result Name	Result LOINC Value
LY27	HLA-B27 Result	26028-1
B27C	Interpretation	69052-9