

Overview

Method Name

ELISA/Chemiluminescence/Indirect Immunofluorescence

NY State Available

Yes

Specimen

Specimen Type

Serum SST

Specimen Required

Collection Container/Tube: Serum gel (red-top tubes are **not acceptable**)

Submission Container/Tube: Plastic vial

Specimen Volume: 2 mL Serum

Collection Instructions:

1. Centrifuge and aliquot 2 mL serum into a plastic vial.
2. Send refrigerated.

Specimen Minimum Volume

Serum: 0.5 mL

Reject Due To

Gross hemolysis	Reject
Gross lipemia	Reject
Gross icterus	Reject

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Serum SST	Refrigerated (preferred)	14 days	
	Ambient	7 days	
	Frozen	30 days	

Clinical & Interpretive

Clinical Information

Refer to www.prometheuslabs.com/about-ibd-precis

Reference Values

An interpretive report will be provided.

Performance**PDF Report**

Referral

Day(s) Performed

Monday through Friday

Report Available

6 to 8 days

Performing Laboratory Location

Prometheus Laboratories, Inc.

Fees & 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 account representative. For assistance, contact [Customer Service](#).

Test Classification

IBD Precis is a laboratory-developed test that was developed, and analytically and clinically validated by Prometheus Laboratories Inc. under federal Clinical Laboratory Improvement Amendments (CLIA) guidelines, and is performed exclusively in our high complexity CLIA certified (05D0917432) and College of American Pathologists (CAP) accredited (6805501) clinical laboratory. As a laboratory developed test, it has not been cleared or approved by the US FDA.

CPT Code Information

81599

LOINC® Information

Test ID	Test Order Name	Order LOINC® Value
FPIBD	IBD Precis	Not Provided
Result ID	Test Result Name	Result LOINC® Value

FPIBD	IBD Precis	Not Provided
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