Overview

Special Instructions
- Anti-Enterocyte Antibody (AEA) Clinical Form

Method Name
Indirect Immunofluorescence

NY State Available
No

Specimen

Specimen Type
Serum Red

Specimen Required
Specimen Type: Serum
Container/Tube: Red
Specimen volume: 1 mL

Collection Instructions: Collect blood in a red-top no additive tube and submit 1 mL of serum shipped frozen.

REQUIRED to accompany all specimens (testing will not proceed until all requirements are met):
1. Completed clinical summary/medical history form
2. See Special Instructions for a copy of the form.

Specimen Minimum Volume
1 mL

Reject Due To

<p>| | | |</p>
<table>
<thead>
<tr>
<th></th>
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<tbody>
<tr>
<td>Hemolysis</td>
<td>NA</td>
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<tr>
<td>Lipemia</td>
<td>NA</td>
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</tr>
<tr>
<td>Icterus</td>
<td>NA</td>
<td></td>
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<tr>
<td>Other</td>
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Specimen Stability Information

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<tr>
<th>Specimen Type</th>
<th>Temperature</th>
<th>Time</th>
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<tr>
<td>Serum Red</td>
<td>Frozen</td>
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Clinical and Interpretive

Reference Values
IgG: Negative
IgA: Negative
IgM: Negative

Performance

PDF Report
No

Day(s) and Time(s) Test Performed
Batched

Analytic Time
4 - 8 weeks

Maximum Laboratory Time
4 - 8 weeks

Performing Laboratory Location
The Children's Hospital of Philadelphia Main Bldg 5th Floor Rm 5203

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

These tests were developed and their performance characteristics determined by the Pathology Department at The Children's Hospital of Philadelphia. They have not been cleared or approved by the U.S. Food And Drug Administration. The FDA has determined that such clearance or approval is not necessary. These tests are used for clinical purposes. It should not be regarded as investigational or for research. This Laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) as qualified to perform high complexity clinical laboratory testing.

CPT Code Information
88346
88350 x 2

LOINC® Information
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<th>Order LOINC Value</th>
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<tr>
<td>FAEAB</td>
<td>Anti-Enterocyte Antibodies</td>
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<table>
<thead>
<tr>
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<th>Result LOINC Value</th>
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