Overview

Method Name
Quantitative Enzyme-Linked Immunosorbent Assay (ELISA)

NY State Available
Yes

Specimen

Specimen Type
Serum

Specimen Required
Draw blood in a plain, red-top tube(s) or serum gel tube(s). Allow serum to clot completely at room temperature. Spin down and send 2 mL serum frozen in a plastic vial.

Specimen Minimum Volume
0.3 mL

Reject Due To

<table>
<thead>
<tr>
<th>Hemolysis:</th>
<th>NA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thawing:</td>
<td>Warm reject; Cold reject</td>
</tr>
<tr>
<td>Lipemia:</td>
<td>NA</td>
</tr>
<tr>
<td>Icterus:</td>
<td>NA</td>
</tr>
<tr>
<td>Other:</td>
<td>NA</td>
</tr>
</tbody>
</table>

Specimen Stability Information

<table>
<thead>
<tr>
<th>Specimen Type</th>
<th>Temperature</th>
<th>Time</th>
<th>Special Container</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum</td>
<td>Frozen (preferred)</td>
<td>120 days</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Refrigerated</td>
<td>120 days</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Ambient</td>
<td>48 hours</td>
<td></td>
</tr>
</tbody>
</table>

Clinical and Interpretive

Reference Values
0.0 - 2.2 ng/mL

SCC antigen levels alone should not be interpreted as evidence of the presence or absence of malignant disease. In patients with known or expected cancer, other tests and procedures must be considered for diagnosis and patient
management. Results obtained with different assay methods or kits cannot be used interchangeably.

Performance

PDF Report
No

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

Analytic Time
1 - 8 days

Maximum Laboratory Time
3 - 10 days

Performing Laboratory Location
ARUP Laboratories

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

Analyte Specific Reagents (ASR) are used in many laboratory tests necessary for standard medical care and generally do require U.S. Food and Drug Administration approval. This test was developed and its performance characteristics determined by ARUP Laboratories, Inc. It has not been approved by the U.S. Food and Drug Administration. This test should not be regarded as investigational or for research use.

CPT Code Information

86316

LOINC® Information

<table>
<thead>
<tr>
<th>Test ID</th>
<th>Test Order Name</th>
<th>Order LOINC Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSCC</td>
<td>Squamous Cell Carcinoma</td>
<td>9679-2</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Result ID</th>
<th>Test Result Name</th>
<th>Result LOINC Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Z2651</td>
<td>Squamous Cell Carcinoma Antibody</td>
<td>9679-2</td>
</tr>
</tbody>
</table>