Overview

Useful For
Assessing calcium states during liver transplantation surgery, cardiopulmonary bypass, or any procedure requiring rapid transfusion of whole blood in neonates and critically ill patients

Second-order test in the evaluation of patients with abnormal calcium values

Method Name
Ion-SelectiveElectrode(ISE)

NY State Available
Yes

Specimen

Specimen Type
Serum SST

Specimen Required
Container/Tube: Serum gel or serum gel microtainer

Specimen Volume: Full tube

Collection Instructions:
1. Allow blood to clot for 30 minutes.
2. Serum gel tube/microtainer must be centrifuged within 1 hour of draw time. Centrifuge with stopper in place for 7 minutes at 3,000 rpm to ensure that the gel barrier separates the serum and cells.

Forms
If not ordering electronically, complete, print, and send a Renal Diagnostics Test Request (T830) with the specimen.

Specimen Minimum Volume
1.75 mL in a 3.5 mL (50% full) in serum gel tube or 1 full serum gel microtainer

Reject Due To

<table>
<thead>
<tr>
<th>Gross hemolysis</th>
<th>Reject</th>
</tr>
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<tbody>
<tr>
<td>Other</td>
<td>Tubes less than 50% full, specimens that have been aliquoted, opened, or poorly centrifuged</td>
</tr>
</tbody>
</table>

Specimen Stability Information
Ionized calcium, which accounts for 50% to 55% of total calcium, is the physiologically active form of calcium.

Low ionized calcium values are often seen in renal disease, critically ill patients, or patients receiving rapid transfusion of citrated whole blood or blood products.

Increased serum ionized calcium concentrations may be seen with primary hyperparathyroidism, ectopic parathyroid hormone-producing tumors, excess intake of vitamin D, or various malignancies.

Nomograms have been used to calculate ionized calcium from total calcium, albumin, and pH values. However, calculated ionized calcium results have proven to be unsatisfactory. A Mayo study of 114 patients found significant differences between ionized and total calcium in 26% of patients.

Reference Values

IONIZED CALCIUM

< or =13 days old: not established

14 days-<1 year: 5.21-5.99 mg/dL

1-<2 years: 5.04-5.84 mg/dL

2-<3 years: 4.87-5.67 mg/dL

3-<24 years: 4.83-5.52 mg/dL

24-< or =97 years: 4.57-5.43 mg/dL

> or =98 years: not established

pH

< or =13 days old: not established

14 days-97 years old: 7.35-7.48

> or =98 years old: not established

For SI unit Reference Values, see [https://www.mayocliniclabs.com/order-tests/si-unit-conversion.html](https://www.mayocliniclabs.com/order-tests/si-unit-conversion.html)

Interpretation

Serum ionized calcium concentrations 50% below normal will result in severely reduced cardiac stroke work. With moderate to severe hypocalcemia, left ventricular function may be profoundly depressed.
Ionized calcium values are higher in children and young adults.

Ionized calcium result has been adjusted to pH 7.40 to account for changes in specimen pH that may occur during transport. Ionized calcium concentration increases approximately 0.2 mg/dL per 0.1 pH unit decrease.

**Cautions**
Proper specimen handling is necessary to ensure accurate results.

**Clinical Reference**

**Performance**

**Method Description**
The pH and ionized calcium sensors in the GEM Premier 3500 Analyzer are based on the principle of ion-selective electrodes; that is, an electrical potential can be established across a membrane that is selectively permeable to a specific ion.(Operator's manual: GEM Premier 3500 Analyzer. Instrumentation Laboratory, Bedford, MA)

**PDF Report**
No

**Day(s) and Time(s) Test Performed**
Monday through Sunday; Continuously

**Analytic Time**
Same day/1 day

**Maximum Laboratory Time**
1 day

**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 modified from the manufacturer’s instructions. Its performance characteristics were determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

**CPT Code Information**
### LOINC® Information

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<th>Test Order Name</th>
<th>Order LOINC Value</th>
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<table>
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<th>Result ID</th>
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