OneStep
Myoglobin Serum
RapiCard™ InstaTest
Cat # 166773-1

FOR THE QUALITATIVE ASSESSMENT OF HUMAN MYOGLOBIN IN HUMAN SERUM

For in vitro Diagnostic Use

INTENDED USE
The Cortez Diagnostics Myoglobin RapidCard™ InstaTest is an immunochromatography based one step in vitro test. It is designed for qualitative determination of myoglobin in human serum specimens. The presence of myoglobin in serum at level 100 ng/ml or higher can be detected in 5 minutes.

SUMMARY AND EXPLANATION
Cortez Diagnostics Myoglobin RapidCard™ InstaTest is a low molecular weight, cytoplasmic serum protein. Due to its low molecular weight, myoglobin is released more rapidly when muscle cells are damaged than other markers. Serum concentration of myoglobin increases above the normal range as early as 1 hour after myocardial infarction, and peak in approximately 4 to 8 hours after onset. Therefore, myoglobin is better suited for the early diagnosis of acute myocardial infarction (AMI).

Cortez Diagnostics Myoglobin RapidCard™ InstaTest is a sandwich immunoassay. When serum sample is added to sample pad, it moves through the conjugate pad and mobilizes gold anti-myoglobin conjugate that is coated on the conjugate pad. The mixture moves along the membrane by capillary action and reacts with anti-myoglobin antibody that is coated on the test region. If myoglobin is present at levels of 100 ng/ml or greater, the result is the formation of a colored band in the test region. If there is
no myoglobin in the sample, the area will remain colorless. The sample continues to move to the control area and forms a pink to purple color, indicating the test is working and the result is valid.

MATERIAL PROVIDED
1. Cortez Diagnostics Myoglobin RapidCard™ InstaTest device

MATERIALS REQUIRED BUT NOT SUPPLIED
1. Serum collection containers.
2. Timer or clock

STORAGE
Store the test device at 2 to 30°C. Do Not Freeze.

PRECAUTIONS
1. For in vitro diagnostic use only.
2. Do not use product beyond the expiration date.
3. Handle all specimens as potentially infectious.

SPECIMEN COLLECTION AND PREPARATION
1. The serum specimen should be collected under standard laboratory conditions
2. Patient samples performed best when tested immediately after collection. If the sample cannot be tested within 24 hours, freeze until the test can be performed. Allow sample to reach room temperature before proceeding.

Sodium azide can be added as a preservative up to 0.1% without effecting the test results.

QUALITY CONTROL
1. The control band is an internal reagent and procedural control. It will appear if the test has been performed correctly and the reagents are reactive.
2. Good Laboratory Practice recommends the daily use of control materials to validate the reliability of the device. Control materials which is not provided with this test kit are commercially available.

PROCEDURE
1. Bring all materials and specimens to room temperature.
2. Remove the test card from the sealed foil pouch.
3. Place the transfer pipette in the specimen and depress the bulb to withdraw a sample.
4. Hold the pipette in a vertical position over the sample well of the test card and deliver 2-3 drops (100-150 µl) of sample into the sample well.
5. Read the result at 10 minutes.

INTERPRETATION OF RESULTS
Positive:
If two colored bands are visible within 10 minutes, the test result is positive and valid.
Note: Specimens containing very low levels of myoglobin may develop two colored bands over 10 minutes.

**Negative:**
If test area has no color band and the control area displays a colored band, the result is negative and valid.

**Invalid result:**
The test result is invalid if a colored band does not form in the control region. The sample must be re-tested, using a new test device.

### LIMITATIONS OF THE PROCEDURE
1. A number of conditions, other than myocardial infarction, including polymyositis, dermatomyositis, systemic lupus erythematosus, shock, severe renal failure, or muscle damage caused by trauma, ischemia and inflammation, can cause elevated levels of myoglobin. These conditions should be considered with appropriate clinical evidence.
2. Recent cardioversion or an anginal episode may increase myoglobin level.
3. Testing 12 hours or later after onset of myocardial infarction can produce misleading results, because serum levels may already have returned to normal range.
4. As with all diagnostic tests, a definitive clinical diagnosis should not be based on the result of a single test, but should only be made by the physician after all clinical and laboratory findings have been evaluated.

### EXPECTED VALUES
Normal serum myoglobin levels range from 30 to 90ng/ml. After 1 hour of the onset of myocardial infarction, serum myoglobin level can elevate to 200ng/ml or even higher. During the peak hour, myoglobin level can be as high as 900 ng/ml. The level of myoglobin usually returns to normal 12 hours after the onset of the myocardial infarction. Elevated myoglobin level has also been observed in patients with other diseases as mentioned in LIMITATIONS OF THE PROCEDURE.

### PERFORMANCE CHARACTERISTICS
**Sensitivity:**
Cortez Diagnostics Myoglobin RapidCard™ InstaTest can detect myoglobin in serum with concentration of 100 ng/ml or greater.

**Accuracy:**

<table>
<thead>
<tr>
<th></th>
<th>Negative (0 ng/ml)</th>
<th>Myo (13 – 66 ng/ml)</th>
<th>Myo (&gt; 100 ng/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of specimen</td>
<td>66</td>
<td>50</td>
<td>100</td>
</tr>
<tr>
<td>Negative</td>
<td>66</td>
<td>34</td>
<td>0</td>
</tr>
<tr>
<td>Positive</td>
<td>0</td>
<td>16</td>
<td>100</td>
</tr>
<tr>
<td>Specificity/Sensitivity</td>
<td>100%</td>
<td>32.0%</td>
<td>100%</td>
</tr>
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**Interference testing:**
The following substances were added to myoglobin negative and 100 ng/ml myoglobin spiked serum samples. No interference was found with any of the substances at the following concentrations:
REFERENCES

<table>
<thead>
<tr>
<th>Date Adopted</th>
<th>Reference No.</th>
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<tr>
<td>2005-09-27</td>
<td>DA-Myoglobin Serum-2008</td>
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