

Intended Use

This CK-MB Isoenzyme Control has been assayed for CK-MB and is intended to be used as a human serum control product for CK-MB determinations. **Rx Only**

Introduction

The determination of the activity of the CK-MB isoenzyme is a valuable diagnostic tool in the evaluation of many pathological conditions, including, acute myocardial infarction. These controls have been designed for use in both manual and automated systems. The use of control material is necessary to estimate test precision in a test system and to detect systematic analytical instrument variation. These control levels are available for the clinical laboratory to evaluate the patient CK-MB isoenzyme level in the normal and abnormal conditions. Since these controls consist of all human preparation, they can be run side by side with the patient sample through all phases of the test method. These human controls eliminate possible altered values found in non-human based materials.

Product Description

These CK-MB isoenzyme controls are prepared from human serum and human derived isoenzyme and are available in two levels. This product is freeze-dried for extended shelf life.

Precautions

1. Check the vial label for the range of isoenzyme values specific for the lot number of control being used.
2. These products have been found to be non-reactive for Hepatitis B Surface Antigen (HBsAg) and negative for antibody to Human Immunodeficiency Virus (HIV), when tested by the FDA approved third generation methods. No methods for HBsAg and HIV can offer total assurance that products derived from human blood will not transmit these diseases. Therefore, human serum products and patients samples should be considered potentially hazardous and handled as if capable of transmitting infectious agents.
3. CAUTION: control material contains azides that can react with copper or lead plumbing to form explosive azides. After use, flush with copious amounts of water to prevent azide build up.

Storage and Stability

CK-MB Isoenzyme Controls are stable until the date indicated on the vial when stored at 2-8°C.

Reconstituted vials are stable for seven days when stored at 2-8°C. If turbid or gross contamination appears after reconstitution, discard immediately.

Procedure

1. Remove vials from refrigerator and allow to warm to room temperature for 15 to 20 minutes.
2. Remove the cap and rubber stopper from vials. Volumetrically add exactly 3.0±0.05ml of distilled or deionized water using a calibrated pipette. The water used for reconstitution should be at room temperature (22-28°C).
3. Recap the vial and gently swirl 10 times.
4. Let the vials remain at room temperature for 20 minutes, then invert gently 10 times.
5. Let the vials remain at room temperature for an additional 20 minutes. Then invert 10 times and gently swirl.
6. Use immediately or refrigerate at 2-8°C for future use.

Limitations

1. These CK-MB controls have been evaluated using the manufacturer's CK-MB reagent.
2. Each lot of control has its own determined value.
3. Individual laboratories may not obtain the mean values as listed for each lot. Technique, equipment and experimental error may produce slightly different values, however, the values should fall within the expected range. Each laboratory should determine their own mean values for this product.










Expected Values





See vial labels for expected ranges.




References

1. Gerhardt W., et al. Clin Chem Acta 78:29 (1977).
2. Giegel J.L., et al. Clin Chem 28:1364 (1982).
3. Mercer D.W., Clin Chem 20:36 (1974).
4. Neumetier D., et al. Clin Chem Acta 73:445 (1976).
5. Roberts R., et al. The Lancet 319 (1977).
6. Vaidya H.C., et al. Clin Chem 32:657 (1986).
7. Wicks R.W., et al. Clin. Chem 28:54 (1982).
8. Willerson J.T., et al. Proc Natl Acad Sci USA 74:1711 (1977).

Symbol Key

 Use by (YYYY-MM-DD)	 Lot and batch code
 Catalog number	 Manufacturer
 <i>In vitro</i> diagnostic medical device	 Temperature limitation
 Consult instructions for use	Rx Only: Prescription Use Only
 CE mark	 Authorized representative in the European Community

 C7562-12  Manufactured for HORIBA Instruments Incorporated - Pointe Brand
5449 Research Drive Canton, MI 48188  2°C-8°C 

 Manufactured for HORIBA Instruments Incorporated – Pointe Brand 5449 Research Drive, Canton, MI 48188	
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Certified to Perform Reagents

The Pointe reagents are certified to be manufactured according to specified parameters. Any Pointe reagent product not meeting specifications through its listed expiration date will be remedied immediately without charge.

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