diatron••

ASO CALIBRATOR

2°C

Packaging: 5 x 1 x 1 mL

INTENDED USE

Calibration material for the preparation of calibration curves for the quantitative determination of Antistreptolysin-O (ASO) using Medicon reagent with Diatron Pictus® P700 and P500 analyzers. For in vitro Diagnostic use only by trained laboratory professionals.

COMPOSITION

Liquid, non-hemolyzed, filtered human plasma with stabilizers and preservative.

- ▲ WARNINGS PRECAUTIONS
 - For in vitro use only
 - Biological materials of human origin contained in this product were tested for Anti-HCV, HbsAg and Anti-HIV 1 and 2 using FDA approved
 methods and were found to be non-reactive. As there is no known test method that can offer complete assurance that products derived from
 human blood will not transmit infectious agents this product should be handled as a potentially infectious material.
 - Any serious incident that may occur in relation to this device must be reported by the user to the manufacturer and the competent authority
 of the country in which the user and/or the patient is established!

▲ PREPARATION

Material ready to use, Swirl the vials gently before use. Avoid foaming. Replace cap immediately after use and store at 2-8°C. Unsuitable storage, handling, or errors during the analytical procedure may give erroneous results.

MATERIALS REQUIRED BUT NOT PROVIDED WITH THE KIT

- ASO reagent kit
- Automated biochemical analyzer
- Common laboratory equipment.

▲ STORAGE - STABILITY

The material unopened remains stable at 2-8°C until the expiry date stated on the label. Once opened the material remains stable for 1 month when stored tightly capped at 2-8°C.

▲ DETERIORATION

The material should not be used:

- · After prolonged exposure to direct sunlight or high temperature
- After the expiry date
- When microbial growth is evident

TEST PROCEDURE

Refer to the appropriate User's Guide and the accompanying Instrument Setting Sheet for analyzer-specific assay instructions for serum/plasma application.

If any trends or sudden shifts in values are detected, review all operating parameters. Control recovery should lie within acceptable intervals. Results outside the specified values even after recalibration could be due to reagent/calibrator/control deterioration, unsuitable storage conditions, instrument malfunction, or error during test procedure.

Every laboratory should establish its own corrective measures in case control recovery lies outside acceptable intervals.

Make sure the LOT on the vial is the same as on the value sheet accompanying the material.

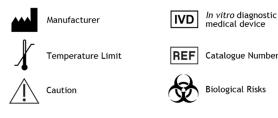
▲ WASTE DISPOSAL

Dispose of all waste materials in accordance with local guidelines. Material safety data sheet is available for professional use upon request.

ASSIGNED VALUES – Lot specific

Please refer to the value sheet for the specific lot available at https://medicondoc.com

SYMBOLS



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