

# **BioMed-PATHOTROL**

REF: (1 Vial) 5 ml REF: (5 Vials) 5 ml each.

### **Description:**

BioMed-PATHOTROL is a lyophilized human-based control serum.

### **Storage and stability:**

Unopened bottles must be stored at 2-8 °C and are Stable up to the expiration date printed on the labels. After reconstitution the serum can be used, stored At 2-8°C and protected from light , in between 5 days.

Bilirubin is stable 1 day, refrigerated. The frozen serum is stable for at least 30 days at  $< -20^{\circ}$ C.

#### **Warnings and Precautions:**

Each individual blood donation used for this serum was found negative when tested with FDA-approved methods on HBsAg, anti-HIV 1+2 and anti-HCV. Nevertheless the serum should be treated for safety reasons always as a potentially infectious material.

### **Preparation:**

Reconstitute carefully by adding exactly 5 ml of Distilled water. Allow the vial to stand for 30 minutes, swirling occasionally. Avoid foaming i.e. do not shake! For alkaline phosphatase the control serum Should stand for 2 hours at + 25 °C before use.

#### **Procedure:**

Refer to the package inserts of the reagent kits

## **Assay Values and Ranges:**

Assay values for analytes for which approved Reference methods are available were determined According to Guidelines of the German Federal Medical Council [Bundesaerztekammer] from 1987 (reference method values) [3].

Ranges of acceptance were calculated as assigned Value  $\pm$  three times the maximum tolerable deviation of a single value according to the Guidelines of the German Federal Medical Council [4].

#### Literature:

1. Röhle G, Siekmann L. Quality assurance of Quantitative determination. In: Thomas L, editor. Clinical laboratory diagnostics. 1st ed. Frankfurt: TH-Books Verlagsgesellschaft; 1998. p. 1393-1401

- 2. Biosafety in Microbiological and Biomedical Laboratories. U.S. Department of Health and Human Services, Washington 1993 (HHS Publication No. [CDC] 93-8395).
- 3. Richtlinien der Bundesärztekammer zur Qualitätssicherung in medizinischen Laboratorien. Deutsches Ärzteblatt 1988;85: B519–B532.
- 4. Richtlinie der Bundesärztekammer zur Qualitätssicherung quantitativer Laboratoriumsmedizinischer Untersuchungen. Deutsches Ärzteblatt 2002; 98: A 2747-59.



