

Art.-Nr.: KG2617
Ch.-B.: 449PD
Inhalt: 6 x 0.5 ml
Verw. bis: 2009-07

PREPARATION

Carefully reconstitute each vial of lyophilised red cell haemolysate base with 0.5 ml of redistilled water. Close bottle and allow to stand for 15 minutes before use. Ensure contents are completely dissolved by swirling gently. Invert bottle to ensure that all traces of dry material are dissolved. Avoid the formation of foam by shaking.

STABILITY

The lyophilised red cell haemolysate base is stable until the expiry date when stored at +2 to +8°C. The G-6-PDH control is stable for 5 days at +2 to +8°C after reconstitution.

Digitonin sample pretreatment assays

After reconstitution the haemolysate requires no further pretreatment.

Concentration = 306 ± 61 U/l 37°C

SAFETY PRECAUTIONS

Human source material from which this product has been derived has been tested at donor level for the Human Immunodeficiency Virus (HIV 1, HIV 2) antibody, Hepatitis B Surface Antigen (HbsAg), and Hepatitis C Virus (HCV) antibody and found to be NON-REACTIVE.

FDA approved methods have been used to conduct these tests.

However, since no method can offer complete assurance as to the absence of infectious agents, this material and all patient samples should be handled as though capable of transmitting infectious diseases and disposed of accordingly.

For in vitro diagnostic use only. Do not pipette by mouth. Exercise the normal precautions required for handling laboratory reagents.

All solutions contain Sodium Azide. Avoid ingestion or contact with skin or mucous membranes. In case of skin contact, flush affected area with copious amounts of water. In case of contact with eyes or if ingested, seek immediate medical attention.

Sodium azide reacts with lead and copper plumbing, to form potentially explosive azides. When disposing of such reagents flush with large volumes of water to prevent azide build up. Exposed metal surfaces should be cleaned with 10 % sodium hydroxide.

18 Nov '08