

μTASWako AFP-L3 Sample Dilution Buffer

Symbols in Product Labeling

REF	Catalog number		Consult instructions for use
IVD	In vitro diagnostics medical device		Use by (last day of the month)
CONT.	Contents of kit		Caution
AC.	Accessory		Manufacturer
Adaptor	Adaptor		Temperature limitation
Holder	Holder		
LOT	Batch code		
EC REP	Authorized Representative in the European Community		
BL	Blank		
CAL	Calibrator		
CONTROL	Control		
CONC.	Assigned value		

Intended use

The μTASWako AFP-L3 Sample Dilution Buffer is designed to be used with μTASWako AFP-L3 for the quantitative determination of AFP-L3% in human serum.

Reagents and ingredients

μTASWako AFP-L3 Sample Dilution Buffer 3 × 10 mL
51 mmol/L Phosphate buffer, pH 6.0

Store at 2 – 10°C. (Do not freeze)

Reagent preparation

Use the Sample Dilution Buffer as supplied. Sample Dilution Buffer is stable until the expiration date printed on the label when stored at 2 – 10°C. (Do not freeze)

Procedure

When the AFP concentration exceeds the reportable range, the sample needs to be diluted by mixing the sample with the Sample Dilution Buffer in an appropriate container. Refer to the μTASWako i30 Instruction Manual for more details about the dilution procedure.

Warnings and precautions

- (1) For in vitro diagnostic use.
- (2) Not to be used internally in human or animals.
- (3) Do not use any buffer or water other than this diluent for sample dilution.
- (4) Do not use the Sample Dilution Buffer for any other purpose than described herein.
- (5) Do not use the Sample Dilution Buffer past the expiration date stated on each container label.
- (6) Do not use the Sample Dilution Buffer in any other procedures than those described herein. Refer to the μTASWako AFP-L3 package insert and the μTASWako i30 Instruction Manual for additional information on the procedure.
- (7) Store the Sample Dilution Buffer under the specified conditions.
- (8) Do not use the Sample Dilution Buffer if they are frozen by mistake, since optimal results may not be obtained with frozen, then thawed Sample Dilution Buffer.
- (9) If the Sample Dilution Buffer comes in contact with mouth, eyes, or skin, wash the exposed area immediately with a large amount of water. Contact a physician if necessary.
- (10) Dispose of Sample Dilution Buffer according to your local or national regulations.
- (11) FDA UDI (Unique Device Identification) is on the box. Keep box until reagent is finished.

This product contains components classified as follows according to the European Regulation :

Hazard designation of product



Mixture containing :

5-Chloro-2-methyl-2H-isothiazol-3-one [EC No 247-500-7] and 2-Methyl-2H-isothiazol-3-one [EC No 220-239-6] (3:1)

Information pertaining to particular dangers for man and environment

Hazard statement

May cause an allergic skin reaction.

Precautionary statement

Avoid breathing vapours/spray.

Contaminated work clothing should not be allowed out of the workplace.

Wear protective gloves/ protective clothing/ eye protection.

If on skin : Wash with plenty of soap and water.

If skin irritation or rash occurs : get medical advice/attention.

Wash contaminated clothing before reuse.

Dispose of contents or container in accordance with local/regional/national/international regulation.

Ordering information

Code No.	Product	Package
997-61501	μTASWako AFP-L3 Sample Dilution Buffer	3 × 10 mL

Revision Date ; July. 2, 2018



μ TASWako i30, its IVD reagent kits and the consumables developed by Wako use Wako's proprietary LBA-EATA assay technologies and microfluidic technologies licensed from Caliper Life Sciences, Inc.

Manufactured by
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