

MICROPROTEIN CONTROL LEVEL I-II

En

REF: VT-038 Microprotein Control

Level I

Microprotein Control

Level II

5 x 2 mL

5 x 2 mL

Changes made in the instructions for use are marked as grey.

Package insert instructions must be carefully followed. Reliability of assay results cannot be guaranteed if there are any deviations from the instructions in this package.

INTENDED USE

REF: VT-039

Validity Microprotein Control Level I-II are for quality control of the Microprotein Liquid assay.

CONTENTS / MATERIALS PROVIDED

Validity Microprotein Control Level I

REF: VT-038

Package: 5 x 2 mL Liquid

Validity Microprotein Control Level II

REF: VT-039

Package: 5 x 2 mL Liquid

For use with;

Validity Microprotein Liquid reagent Validity Microprotein Calibrator

VT-038 / VT-039 contain the human serum. Sodium azide (0.09%) is added as preservative.

Materials Required But Not Provided:

- 1. Class A volumetric pipette for liquid transfer
- Distilled or deionized water meeting the specifications equivalent to USP (United States Pharmacopeial Convention) purified water.

CONTROL STABILITY

Temperature-Conditions	Stability
Unopened at +2/+8°C	Expiry date on the vial.
Opened and stored at +2/+8°C	30 days

PREPARATION OF CONTROL

Control is ready for use.

Rev: V1.0 Date: 12.2023

INDICATIONS OF INSTABILITY OR DETERIORATION

Presence of extreme turbidity or microbial growth may indicate deterioration.

PRECAUTIONS



Human source material. Treat as potentially infectious material. Each plasma donor used in the preparation of this product has been tested by an FDA-

approved method and found negative for the presence of HIV 1/2 HBsAg, HCV, HIV-Ag antibodies. However, none of the known testing methods can offer complete assurance that the hepatitis B virus, Human Immunodeficiency Virus (HIV) or infectious agents are not present. All human-based products should be handled in accordance with Good Laboratory Practice (GLP) principles using appropriate precautions. Safety data sheets are available at www.validity.com.tr or you may contact your local representative.

WARNINGS

IVD: For in Vitro Diagnostic use only.

Do not use expired reagents.

Reagents with two different lot numbers should not be interchanged.

For professional use.

Follow Good Laboratory Practice (GLP) guidelines.

Contains sodium azide.

CAUTION: Human source samples are processed with this product. All human source samples must be treated as potentially infectious materials and must be handled in accordance with OSHA standards.

Danger

H317 :May cause allergic skin reaction.

Precaution

P280 :Use protective gloves / clothes /

glasses / mask.

P264 :Wash your hands properly after

using.

P272 :Contaminated work clothes should

not be allowed to be used outside of

the workplace.

Intervention

P302+P352 :Wash with plenty of water and soap if

it contacts with skin.

P333+P313 :Seek medical help if it irritates your

skin or develops rash.

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P362+P364 :Remove contaminated clothes and

wash properly before using.

Disposal

P501 :Dispose the vials and contents

according to the local regulations.

REFERENCES

 Council Directive (2000/54/EC). Official Journal of the European Communities No. L262 from Oct. 17, 2000.

- EU-Dir 1999/11 Commission Directive of 8 March 1999 adapting to technical progress the principles of Good Laboratory Practice as specified in Council Directive 87/18/EEC
- Clinical and Laboratory Standards Institute, H26-A2, Validation, verification, and quality assurance of automated hematology analyzers; Approved Guideline - Second Edition.
- 4. Gabbay, K.H., Hasty, K., Breslow, J.L., Ellison, R.C., Bunn, H.F., and Gallop, P.M., J. Clin. Endocrinol. Metab. 44, 859 (1977).
- 5. US Department of Labor, Occupational Safety and Health Administration, 29 CFR Part 1910.1030, Occupational Exposure to Bloodborne Pathogens.



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