

ARCHEM RHEUMATOID CONTROL AND CALIBRATOR

En

REF: TA111S REF: TA121S ASO Standard Calibrator RF Standard Calibrator

5x1 mL 5x2 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

Archem Rheumatoid Control and Calibrator is for quality control and calibration of the parameters stated in the value sheet.

CONTENTS / MATERIALS PROVIDED

ASO Standard (Calibrator)
REF: TA111S

Package: 5 x 1 mL

RF Standard (Calibrator)

REF: TA121S Package: 5 x 2 mL

Archem Rheumatoid Control and ASO, CRP, RF Calibrator 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.

CALIBRATOR AND CONTROL STABILITY

Rev: V1.1 Date: 09.2021

Temperature-Conditions	Stability
Powder form, stored in dark place at	Expiry date on the
+2/+8 °C	vial
Diluted and stored in dark place at +25°C,	8 hours
Diluted and stored in dark place at +2/+8°C	2 days
Diluted, stored in dark place and frozen at -20°C	180 days

LIMITATIONS

- Please open the vial caps carefully. When you open, be careful not to scatter any powdery substance around or to escape from the vial.
- Dissolve with distilled water with volume stated on the vial. Injector should not be used for the transfer process since there may be errors between 5-20% in liquid transfer with the injector. Use calibrated micropipettes.
- Temperature of dry serum in the vial and distilled water must be +20/+25°C. After adding distilled water, close the vial cap tightly and store at +25°C around 5-10 minutes.
- 4. Wait for 30 minutes for dissolving process and mix thoroughly by gently inverting the vial at regular intervals, do not shake. Avoid formation of bubbles or foam. Protect from light. It is recommended to use a rotational mixer for routine mixing procedures.
- 5. After reconstitution, the control and calibrator serums are usually divided into small quantities (150-250 microliters) into Eppendorf tubes or sample cups of the device and stored in the refrigerator for freezing process. For serums prepared in this way, it is absolutely necessary to leave the serum at +25°C for 30 minutes before dividing it into small quantities. Do not refreeze after the serum is frozen and thawed once.
- 6. Control and calibrator serum precipitation is faster than normal serum. In order for the first and last parts to be homogeneous and to avoid precipitation, perform the process as fast as possible during separation.
- 7. The quality of the distilled water to be used in the dilution of the calibrator serum is very important. There may be significant deviations in the values due to bacterial contamination.
- 8. It is necessary to be careful against infectious agents in calibrator serum measurements.

PREPARATION OF CALIBRATOR

Lyophilized control and calibrator serum should be reconstituted by adding distilled or deionized water with the amount stated on the label. Close the vial and wait for 20 minutes. Dissolve the contents of the vial by swirling gently to avoid the formation of foam. Do not shake.

INDICATIONS OF INSTABILITY OR DETERIORATION

Presence of extreme turbidity or microbial growth may indicate deterioration.



ARCHEM RHEUMATOID CONTROL AND CALIBRATOR

En

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.archem.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.

REFERENCES

- Burtis CA, Ashwood ER, Bruns DE, editors. Tiets Textbook of Clinical Chemistry and Molecular Diagnostics, 4th ed. St. Louis, MO, Elsevier Saunders; 2006:2263.
- S.Dean Allison, Mark C.Manning, Theodore W.Randolph, Kim Middleton, Ashley Davis, John
 F.Carpenter. Optimization Of Storage Of Lyophilized Actin Using Combinations Of Disaccharides And Dextran. Journal Of Pharmaceutical Sciences.89/2,199-214(2000)

TRADEMARKS

Archem Rheumatoid Control and Calibrator is a trademark of ARCHEM Sağlık Sanayi ve Tic. A.Ş. in various jurisdictions.



Archem Sağlık Sanayi ve Tic. A.Ş.

Mahmutbey Mah. Halkalı Cad. No:124 Kat:4 Bağcılar/İstanbul/Turkey

Tlf: + 90 212 444 08 92 Fax: +90 212 629 98 89

info@archem.com.tr www.archem.com.tr



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.

P362+P364 :Remove contaminated clothes and wash

properly before using.

Disposal

Rev: V1.1

P501 :Dispose the vials and contents according to

the local regulations.

Date: 09.2021



ARCHEM RHEUMATOID CONTROL AND CALIBRATOR





