

Order references

Reagents

REF		CONT
ASLTX-B00	Universal kit	1 x 50 ml R1 + 1 x 5 ml R2
ASLTX-H00	Universal kit	2 x 70 ml R1 + 1 x 15 ml R2

Other necessary products

REF		CONT
ASREH-001	Anti-Streptolysin O High Calibrator	1 x 1 ml
ASREH-005	Anti-Streptolysin O High Calibrator	1 x 5 ml
ASCON-002	Anti-Streptolysin O Control	1 x 2 ml

Field of application - Purpose

In vitro diagnostic reagent for the quantitative determination of antistreptolysin O in samples of human origin by immunoturbidimetry on photometric systems.

Medical benefit - Scientific validity

In 1928, Todd demonstrated the production by group A streptococci of a lysine from human red blood cells. He demonstrated its antigenic character, which causes the production of circulating antistreptolysin O (ASO) antibodies after infection by the microorganism. Streptolysin O is a soluble protein of 69 kDa that binds to the erythrocyte membrane and permeabilises it. The assay of antistreptolysin O enables monitoring the progression of secondary complications related to this condition, which can be acute rheumatic fever or acute glomerulonephritis.

Method principle

The latex particles are stabilized in colloidal form with anti-antistreptolysin O antibodies specifically directed against the antistreptolysin O. The reaction of these particles with the antistreptolysin O, present in a biological sample, causes specific agglutination of the latex particles. This agglutination, directly proportional to the antistreptolysin O concentration of the sample, is read at 600 nm.

Warning and precautions

- For in vitro diagnostic use only.
- Must be handled by qualified personnel under the responsibility of a biologist.
- The human-origin products have been screened and found negative for HIV 1 and 2 antibodies, HCV antibodies and HBAG, but they must nevertheless be handled as potentially infectious products.
- These products contain sodium azide. Products containing sodium azide must be handled with care: avoid ingestion and contact with the skin or mucous membranes.
- Sodium azide becomes explosive on contact with heavy metals such as copper or lead.

Samples

Collection conditions

Collect specimens using standard laboratory techniques; use only suitable procedures, tubes or collection containers.

Sample type

Non-haemolysed and non-lipaemic serum and plasma (heparin or EDTA).

Storage and stability of specimens

Temperature	Stability
- 20 °C	≤ 6 months
4 - 8 °C	≤ 8 days
20 - 25 °C	≤ 2 days

This information comes from data originating from “Tietz Clinical Guide to Laboratory Tests” and from “WHO”.

Reagents

Composition and concentrations/Storage

Active ingredients:

Reagent R1: none.

Reagent R2: Suspension of latex colloidal particles covered with streptolysin O.

Other ingredients:

Reagent R1: buffer, inorganic salt and preservative.

Reagent R2: buffer, polystyrene, inorganic salt and preservative.

Storage temperature:

Reagent R1: 2 - 8°C.

Reagent R2: 2 - 8°C

Preparation

Ready to use.

Storage and stability

Reagents are stable until the expiration date printed on the packaging (months passed), under the following recommended storage and handling conditions:

- Unopened vial stored at temperature indicated on packaging.
- Opened vial: closed immediately after use or placed on closed analyser intended for this purpose, not contaminated by handling and stored at the temperature indicated on the packaging.

Note:

- Do not freeze the reagents.
- Nanoparticle-based reagents can settle over time. It may be necessary to delicately mix by repeated turning.

Other materials required

Usual laboratory equipment including an analytical system equipped with a photometric detector.

Calibration

Calibration

The calibration curve is performed by using the calibration kit indicated in the “Order references” section. The zero point of the calibration curve is performed with physiological saline solution.

Traceability

The method has been standardised with a benchmark method traceable to the international standard as described in the associated calibrators data sheet (see the “Order references” section).

Calibrate the method when the reagent batch number changes or in case of change in performance (contact the manufacturer if the changes persist) or if quality control requires it.

Quality control

The frequency of controls and the confidence limits must be adapted to the laboratory requirements. The results must be within the defined confidence limits. Each laboratory shall establish corrective measures to be taken if results fall outside the defined limits. Comply with current legislation in the country and local guidelines relating to quality control.

The calibration curve and its stability can be validated using the control materials indicated in the “Order references” section.

Reference values

	Reference values
< 5 years	< 100 IU/mL
Adults	< 200 IU/mL

International units: IU/ml

Conventional units: /

This information coming from data originating from “Clinical guide to laboratory tests”. Each laboratory must check the validity of its values and if necessary establish its own reference values, depending on the population examined.

Analytical performances

The analytical performance data below are given as an indication. The results obtained in the laboratory may differ from these. The analytical performances were determined following the indications of the “Guide technique d’accreditation de vérification (Portée A)/validation (Portée B) des méthodes en biologie médicale”; document SH GTA 04 Révision 01.

Measurement range

19,992 - 864,85 IU/mL

The measurement range is bounded by the quantification and linearity limits. Samples having a concentration greater than the upper limit must be diluted.

Limit of detection

9,231 IU/mL

It is the smallest signal expressed as a quantity or concentration that can be distinguished with a given probability from a reagent blank performed in the same conditions.

The evaluation of the limit of detection is based on the statistical analysis of the observed signal differences between the blanks and samples.

Interferences (Analytical specificity)

There is no known cross-reactivity of the antiserum cited or the antibodies used.

The abnormally coloured and particle-containing samples can cause, depending on the analytical system, assay errors. These samples must be clarified chemically or physically before their assay.

Precision

The precision is evaluated using the repeatability (CV within-run) and reproducibility (CV within-calibration).

	Repeatability (n=30)		Reproducibility (n=30)	
	Average (IU/ml)	CV (%)	Average (IU/ml)	CV (%)
Level 1	105,37	0,76	110,71	3,07
Level 2	176,43	2,35	183,07	2,54
Level 3	229,86	1,46	371,90	3,50
Level 4	358,82	1,86		

Trueness - Accuracy

Trueness, quantified by the bias, is estimated by comparing the mean obtained in the intermediate precision study, based on internal quality control samples, with the expected target value equated to the “true” value of the tested sample.

Accuracy is defined as the closeness of agreement between a measured value and a true value of a measurand (quantity to be measured).

DiAgam allows a bias of 5% compared to the international standard or compared to a reference method traceable to the international standard when it exists.

Limitations of the method

The results of this test should always be interpreted in relation to the patient’s medical history, clinical signs and other findings.

Prozone

By limiting the linearity to the value of the upper limit of the measurement range, no excess antigen effect was observed for samples with a concentration up to 1500 IU/mL.

Matrix effect

The inter-laboratory control samples and controls can yield different results from those obtained with other assay methods because of a matrix effect. In this case, an analysis of the results according to specific target values of the method utilised may be necessary. If in doubt, contact the manufacturer.

Utilisation procedure

Validated automatic applications for different analyzers are available from DiAgam. The utilisation procedure indicated below enables deriving a manual or automatic application of the reagent (NB - comply with the sample/R1/R2 ratios correctly). Please contact the manufacturer for more information.

Mix 2 µl of sample with 235 µl of reagent R1 and incubate the mixture for 5 minutes at 37°C. Then add 30 µl of reagent R2 to the reaction mix. Then read the optical density at a wavelength of 600 nm (OD1) . Incubate at 37°C for 5 min. Perform a second OD measurement at 600 nm (OD2).

This operation must be made with a “reagent blank” sample (physiological saline solution, considered as point zero of the calibration curve), with the calibrators indicated in the “Order references” section and to finish with the samples of unknown concentrations.

The final OD is finally calculated as shown in the following equation:

$$OD_{final} = OD2 - f \times OD1$$

Where f is a factor taking account of the difference in volume between the 2 measurements of OD.






The final OD of the “reagent blank” sample as well as the known calibrator concentrations allows a calibration curve to be drawn. The transfer of the OD measured for an unknown sample on this calibration curve enables its concentration to be determined.


Literature

1. Tietz Textbook of Clinical chemistry and molecular Diagnostics, fourth edition, edited by Carl A. Burtis, Edward R. Ashwood, David E. Bruns, 2006
2. Use of Anticoagulants in Diagnostic Laboratory Investigations & Stability of blood, plasma and serum samples. Publication WHO/DIL/LAB/99.1 Rev. 2. Jan. 2002.
3. Clinical guide to laboratory tests, second edition, edited by Norbert W. Tietz, 1990
4. CLSI. Procedures for the Collection of Diagnostic Blood Specimens by Venipuncture; Approved Standard-Sixth Edition. CLSI document H3-A6 (ISBN 1-56238-650-6). CLSI, 940 West Valley Road, Suite 1400, Wayne, PA 19087-1898 USA; 2007.
5. NCCLS. Procedures and Devices for the Collection of Diagnostic Capillary Blood Specimens; Approved Standard-Fifth Edition. NCCLS document H4-A5 [ISBN 1-56238-538-0]. CLSI, 940 West Valley Road, Suite 1400, Wayne, PA 19087-1898 USA, 2004.

Symbols legend

The following symbols may appear on the packaging and the label:

LOT	Batch code	BUF	Buffer
	Use until	CAL	Calibrator
	Manufacturer	H	High
IVD	In vitro diagnostic medical device	M	Moderate
	Temperature (Storage at)	L	Low
REF	Catalogue reference	4 LEV	4 levels
	Read the usage instructions	5 LEV	5 levels
REAG	Reagent	6 LEV	6 levels
KIT	Kit	CONTROL	Control
CONT	Content		This product meets the requirements of European Directive 98/79 EC concerning in vitro diagnostic medical devices
Ab	Antibody or Antisera		

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