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HBS-PEG-XG

Polyethylene Glycol Solution

Intended Use

HBS-PEG-XG is intended for use as a potentiator in antibody detection, antibody identification and compatibility test procedures.

Summary and Explanation

Polyethylene glycol (PEG-XG) can be used as a potentiator in the antiglobulin test. PEG enhances antibody uptake by excluding water molecules in the diluent. This factor enables a closer proximity of antigen/antibody, resulting in increased antibody binding for a more sensitive test. HBS-PEG-XG can be used in combination with Anti-Human Globulin (Anti-IgG) reagent for compatibility testing, antibody screening and identification.

Principle of the Procedure

The sensitivity of the indirect antiglobulin procedure for the detection and identification of clinically significant antibodies is enhanced by the addition of polyethylene glycol. In addition, a substantial reduction in the incubation time for antigen/antibody mixtures can be achieved when the red cells and serum are suspended in a low ionic strength test environment. HBS- PEG-XG is prepared in a low ionic strength solution base material. HBS-PEG-XG can also be used as an additive when testing eluates, with significantly improved sensitivity when compared to usual methods [3].

Reagent Description

HBS-PEG-XG is a 20% solution of polyethylene glycol 3350 in a low ionic strength medium containing Glycine that creates a low ionic strength environment with sodium azide as a preservative.

Precautions:

1. This reagent contains 0.1% (w/v) sodium azide which is below the national and international regulatory thresholds and when used under normal condition is not chemically hazardous. If this reagent is discarded in the sink, flush with large volumes of water to prevent the buildup of azide.
2. Do not use if the reagent is turbid.
3. The packaging of this product contains dry natural rubber.
4. This reagent is for in vitro diagnostic use only.

Storage

This reagent should be stored at 2-25°C. Do not freeze or expose to elevated temperatures.

Specimen Collection

Blood should be drawn by aseptic methods and the serum or plasma should be tested as soon as possible. If testing is delayed, the samples should be stored at 2-8°C.

Procedure:

Materials Provided

Hemo bioscience PEG-XG

Materials Required but Not Provided

Test tubes (12 x 75 or 10 x 75 mm)
Transfer pipettes
Centrifuge (1000 rcf)
Isotonic or phosphate buffered saline
Timer
Anti-Human Globulin reagent containing IgG
Antiglobulin Control Cells (cells sensitized with IgG)
37°C incubator

Recommended Technique:

1. Donor/Patient red cells should be washed once in isotonic or Phosphate Buffered Saline (PBS) making a final suspension of 2-4%.
NOTE: Reagent red blood cells may be used directly from the vial or in accordance with manufacturer's directions.
2. Add two drops of serum, plasma, or eluate to a test tube.
3. Add one drop of the red blood cell suspension.
4. Add two drops of HBS-PEG-XG and mix thoroughly.
5. Incubate for 10-15 minutes at 37°C in a water bath or heat block. **NOTE: DO NOT centrifuge the test mixture directly**

after the incubation phase. See Performance Limitations section for further information.

6. Examine the tubes for gross hemolysis.
7. Perform AHG test as per manufacturer's instructions.
8. Following centrifugation, all tests should be read immediately and results should be interpreted without delay. Delays may result in disassociation of antigen-antibody complexes leading to falsely negative, or at most, weakly positive reactions.

Quality Control

Quality control of reagents is essential and should be performed each day of use always follow local, state and federal regulations.

Results:

Positive Test

Agglutination or hemolysis of the red blood cells.

Negative Test

No agglutination and no hemolysis of the red blood cells.

Limitations:

1. Polyethylene glycol causes the red blood cells to aggregate, which makes examination for direct agglutination difficult. As a result, the test should **only** be read at the antiglobulin phase.
2. IgM antibodies may not be detected when using the polyethylene glycol procedure, care should be taken to ensure ABO compatibility when cross matching.
3. Polyethylene glycol may precipitate serum globulins. It is imperative that the red blood cells are resuspended thoroughly between washes. The Anti-Human Globulin reagent may be neutralized if cells are not washed thoroughly, resulting in false negative results. It may also be necessary to wash the red blood cells more than four times to remove all unbound human protein.
4. Confirm Anti-Human globulin reagents have not been inactivated using Antiglobulin control cells for negative Indirect Antiglobulin tests.








Specific Performance Characteristics

Each lot of HBS-PEG-XG is tested to ensure appropriate performance. For Technical support, contact Hemo bioscience at 1-866-332-2835.

Bibliography:

1. Nance SJ, Garratty G. Polyethylene glycol: A new potentiator of red blood cell antigen–antibody reactions. *Am J Clin Pathol* 1987; 87:633-5.
2. Löw B, Messeter L. Antiglobulin test in low ionic strength salt solution for rapid antibody screening and crossmatching. *Vox Sang* 1974; 26:53-61.
3. Combs MR, Telen MJ. Testing eluates in Polyethylene glycol (PEG): A sensitive technique for detecting early alloimmunization. [Abstract] *Transfusion* 1989; Supplement 58S.
4. Fung, MK (ed): Technical Manual, 18th ed. AABB, Bethesda MD, 2014.

Glossary of Symbols:

Symbol	Definition
	Batch code
	Manufacturer
	Temperature limitation
	Consult instructions for use.
	Use by YYYY-MM-DD
	For <i>in vitro</i> diagnostic use
	Caution, consult documents.