BLOOD GROUPING REAGENT

Anti-Jka (Monoclonal)  Anti-Jkb (Monoclonal)

Gamma-clone®

By Tube Test

Preservative: <0.1% Sodium Azide 3°C to 10°C  Meets FDA Potency Requirements  IVD

CAUTION: THE PACKAGING OF THIS PRODUCT (DROPPER BULBS) MAY CONTAIN DRY NATURAL RUBBER. DO NOT PIPETTE THIS PRODUCT BY MOUTH, AS THE ABSENCE OF MURINE VIRUS HAS NOT BEEN DETERMINED. DO NOT USE IF MARKEDLY TURBID.

Intended Use:
Gamma-clone Anti-Jka (Monoclonal) and Anti-Jkb (Monoclonal) Blood Grouping Reagents are intended for the detection of the Jk(a+) and Jk(b+) antigens, respectively, on red blood cells by tube test.

Summary of the Test:
Anti-Jka was first reported by Allen, Diamond and Niedziela in 1951, and the first example of the anti-JkB antigen was identified in 1953 by Piault and her co-workers. Both antibodies have been implicated as the cause of hemolytic disease of the newborn and hemolytic transfusion reactions. The phenotype Jk(a−b−), first reported by Pinkerton and associates in a person of Filipino/Chinese descent, is rare among whites, has not been described among African-Americans in the US population, but appears to be relatively common among certain Pacific Island and Asian populations. Immunized individuals of this phenotype may produce an antibody, anti-Jk3, that reacts with all red blood cells that are either Jk(a+) or Jk(b+).

Gamma-clone Anti-Jka (Monoclonal) and Gamma-clone Anti-Jkb (Monoclonal) Blood Grouping Reagents are used to detect the presence of the Jk(a+) and Jk(b+) antigens on donor or patient red blood cells. Typing of donor red blood cells facilitates the selection of antigen-negative units for transfusion to patients with the corresponding antibody. Red blood cell typing also serves as final verification of the identification of an alloantibody in patient or donor serum.

Principle of the Test:
The presence of the Jk(a) and Jk(b) antigens is determined by testing with Anti-Jka and Anti-Jkb by the tube test technique. Agglutination of the test red blood cells constituting a positive test result and indicates the presence of the relevant antigen. No agglutination constitutes a negative test result and indicates that the antigen is not present.

Reagents:
Gamma-clone Anti-Jka (Monoclonal) Blood Grouping Reagent is prepared from IgM antibodies from the human/murine heterohybridoma cell line MS-15 grown in fluid culture and suitably diluted in a proprietary diluent containing bovine albumin to achieve the appropriate level of potency for the test procedure as described. Sodium azide is added as a preservative (at less than 0.1% w/v). Ready for use as supplied.

Gamma-clone Anti-Jkb (Monoclonal) Blood Grouping Reagent is prepared from IgM antibodies from the human/murine heterohybridoma cell line MS-8 grown in fluid culture and suitably diluted in a proprietary diluent containing bovine albumin to achieve the appropriate level of potency for the test procedure as described. Sodium azide is added as a preservative (at less than 0.1% w/v). Ready for use as supplied.

Any Bovine Albumin used in the manufacture of this product is sourced from donor animals of United States origin that have been inspected and certified by US Veterinary Service.

Additional Materials Required:
Gamma-clone Anti-Jka (Monoclonal) or Anti-Jkb (Monoclonal)

Additional Materials Required:
1. Test tubes (12x75 mm or 10x75 mm)
2. Pipettes
3. Isotonic saline or phosphate-buffered (approximately 15 mM) isotonic saline pH 6.5-7.5
4. Centrifuge*
5. An optical aid such as a hand lens or concave mirror
6. Red blood cells of known Kidd phenotypes for use as controls.

*It is the user’s responsibility to validate an accessory device (either listed or otherwise) for its intended use. Validation results should be maintained as part of the laboratory’s records for review by regulatory agencies.

Test Method:
Since this test method applies to either reagent, extreme care should be exercised in selecting and using the appropriate reagent.

1. Place one (1) drop of Gamma-clone Anti-Jka (Monoclonal) or Gamma-clone Anti-Jkb (Monoclonal) into a properly labeled test tube.
2. Add one (1) drop of an approximate 2-5% suspension of the red blood cells to be tested to the test tube (from step 1 above). The red blood cells to be tested should previously have been washed at least one time and resuspended in saline.
3. Mix the test tube contents well by gently shaking the tube and incubate the tube for five (5) to fifteen (15) minutes at room temperature (15°C to 30°C). Incubating for the upper end of the time range may enhance reactivity.
4. Centrifuge the test tube.*
5. After centrifugation, immediately resuspend the red blood cells by gently shaking the test tube and examine for macroscopic agglutination. Negative reactions may be examined with an optical aid; however, microscopic reading is not recommended. Record the results.

Stability of Reaction:
Following centrifugation, the tube test should be read immediately and interpreted without delay.
Quality Control:
The reactivity of blood grouping reagents should be confirmed on each day of use by testing with red blood cells known to be negative and positive for the relevant antigens. Jk(a+b+) red blood cells are the most suitable positive control red blood cells for both Gamma-clone Anti-Jk\(^a\) (Monoclonal) and Gamma-clone Anti-Jk\(^b\) (Monoclonal). Each reagent is satisfactory for use if it reacts only with antigen-positive red blood cells. Immucor Reagent Red Blood Cells are a convenient source of control cells and may be used as supplied.

Interpretation of Results:
Agglutination of the red blood cells constitutes a positive test result and indicates the presence of the relevant antigen.

No agglutination constitutes a negative test result, and indicates the absence of the relevant antigen.

The reaction patterns possible with Anti-Jk\(^a\) and Anti-Jk\(^b\) are shown in Table 1, together with the frequencies of the resulting phenotypes in some ethnic populations.

<table>
<thead>
<tr>
<th>Reagent</th>
<th>Phenotype</th>
<th>Frequency (%)</th>
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<tbody>
<tr>
<td></td>
<td></td>
<td>Caucasians</td>
</tr>
<tr>
<td>Anti-Jk(^a)</td>
<td>-</td>
<td>+</td>
</tr>
<tr>
<td>Anti-Jk(^b)</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Anti-Jk(^b)</td>
<td>0</td>
<td>+</td>
</tr>
<tr>
<td>Anti-Jk(^a)</td>
<td>0</td>
<td>0</td>
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</tbody>
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Table 1: The reaction patterns of Anti-Jk\(^a\) and Anti-Jk\(^b\) and the approximate frequencies of the resulting phenotypes in some ethnic populations.

Limitations:
1. Factors that may cause false test results include the following:
   a. Bacterial or chemical contamination of blood specimens, reagent and/or supplementary materials.
   b. Improper storage of materials.
   c. Aged or stored blood specimens. Such specimens may yield weaker reactions than those obtained with fresh red blood cells.
   d. Too heavy a red blood cell suspension of the specimen.
   e. Improper centrifugation. Proper centrifuge calibration is particularly important to the proper performance of the test. Excessive centrifugation may lead to difficulty in resuspending the red blood cell button in the tube test leading to a possible false positive result. At the same time, inadequate centrifugation may yield unclear red blood cell button patterns and agglutinates that are too readily dispersed leading to a possible false negative result.
   f. Improper examination for agglutination (usually too vigorous shaking). The resuspension of reactions in the tube test procedure must be carried out by gentle shaking. Shaking too vigorously may cause agglutinates to be dispersed.
   g. Deviation from the recommended test procedure such as the omission of test reagents.
2. Positive reactions of red blood cells from persons of unusual Kidd genotypes may be weaker than those reactions obtained with randomly selected positive control red blood cells tested in parallel. For these reasons, caution should be exercised when assigning genetic significance on the basis of test results.
3. Red blood cells that have been enzyme-treated must not be used for testing as either red blood cells under investigation or as a source of control red blood cells because use of these enzyme-treated red blood cells may yield erroneous results.

Specific Performance Characteristics:
Gamma-clone Anti-Jk\(^a\) (Monoclonal) and Anti-Jk\(^b\) (Monoclonal) meets FDA potency requirements. Each lot is tested by insert methods against a panel of antigen-positive and antigen-negative red blood cells to ensure suitable reactivity and specificity. The specificity of the murine monoclonal antibodies secreted by the cell lines used to manufacture these Blood Grouping Reagents has been determined by testing with red blood cells of varying phenotypes.

The performance of this product is dependent upon adhering to the package insert recommended methodology.

For additional information or for technical support, contact Immucor at 855-IMMUCOR (466-8267).

Bibliography: