SRBC IgG (Mouse) ELISA Kit

Catalog Number KA2462
96 assays
Version: 04

Intended for research use only
# Table of Contents

**Introduction** ................................................................................................................................. 3  
  Intended Use .................................................................................................................................. 3  
  Background ..................................................................................................................................... 3  
  Principle of the Assay ....................................................................................................................... 3  

**General Information** ...................................................................................................................... 4  
  Materials Supplied ............................................................................................................................ 4  
  Storage Instruction .............................................................................................................................. 4  
  Materials Required but Not Supplied .................................................................................................. 4  
  Precautions for Use ............................................................................................................................ 5  

**Assay Protocol** ............................................................................................................................... 6  
  Reagent Preparation .......................................................................................................................... 6  
  Sample Preparation ........................................................................................................................... 6  
  Assay Procedure ............................................................................................................................... 6  

**Data Analysis** ................................................................................................................................. 8  
  Calculation of Results ....................................................................................................................... 8  

**Resources** ..................................................................................................................................... 9  
  Reference ......................................................................................................................................... 9  
  Plate Layout ..................................................................................................................................... 10
Introduction

Intended Use

ELISA for the Quantitative Determination of Mouse Anti-Sheep Red Blood Cell (SRBC) IgG in Serum and Plasma.

Background

Recent studies have demonstrated that suppression of anti-SRBC (sheep red blood cell) IgG levels by therapeutic agents serves as a useful indicator of immunosupression.\textsuperscript{1,2} This ELISA allows rapid and quantitative measurement of mouse anti-SRBC IgG levels in serum or plasma samples.

Principle of the Assay

The mouse anti-SRBC IgG ELISA is based on a solid phase enzyme-linked immunosorbent assay (ELISA). The assay uses detergent solubilized SRBC ghosts\textsuperscript{2} for solid phase (microtiter wells) immobilization and horseradish peroxidase (HRP) conjugated anti-mouse IgG antibodies for detection. Test serum or plasma samples are diluted and incubated in the microtiter wells for 45 minutes. The microtiter wells are subsequently washed and HRP conjugate is added and incubated for 45 minutes. Anti-SRBC IgG molecules are thus sandwiched between immobilized SRBC antigens and the detection antibody conjugate. The wells are then washed to remove unbound HRP-labeled antibodies and TMB Reagent is added and incubated for 20 minutes at room temperature. This results in the development of a blue color. Color development is stopped by the addition of Stop Solution, changing the color to yellow, and optical density is measured spectrophotometrically at 450 nm. The concentration of anti-SRBC IgG is proportional to the optical density of the test sample.
General Information

Materials Supplied

List of component

<table>
<thead>
<tr>
<th>Component</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>SRBC coated 96-well Plate (provided as 12 strips of 8 wells)</td>
<td>96 (8 x12) wells</td>
</tr>
<tr>
<td>Enzyme conjugate reagent</td>
<td>11 mL</td>
</tr>
<tr>
<td>Reference standard stock* (Lyophilized)</td>
<td>1 vial</td>
</tr>
<tr>
<td>Diluent</td>
<td>30 mL</td>
</tr>
<tr>
<td>20x Wash Solution</td>
<td>50 mL</td>
</tr>
<tr>
<td>TMB Reagent (One-step)</td>
<td>11 mL</td>
</tr>
<tr>
<td>Stop Solution (1N HCl)</td>
<td>11 mL</td>
</tr>
</tbody>
</table>

*The levels of mouse anti-SRBC IgG are measured in nominal units and are calibrated with reference mouse anti-SRBC serum at Abnova.

Storage Instruction

The kit should be stored at 2-8°C and the microtiter plate should be kept in a sealed bag with desiccant to minimize exposure to damp air. Test kits will remain stable for six months from the date of purchase provided that the components are stored as described above.

Materials Required but Not Supplied

- Precision pipettes and tips
- Distilled or deionized water
- Polypropylene or glass tubes
- Vortex mixer
- Absorbent paper or paper towels
- Micro-Plate incubator/shaker with an approximate mixing speed of ~150 rpm
- Plate washer
- Plate reader with an optical density range of 0-4 at 450 nm
- Graph paper (PC graphing software is optional)
Precautions for Use

- General Instructions
  - Please read and understand the instructions thoroughly before using the kit.
  - All reagents should be allowed to reach room temperature (18-25°C) before use.
  - The assay was designed for use with serum or plasma obtained from mice fourteen days after immunization with SRBC, at which point the immune response originates almost exclusively from IgG.
  - Serum or plasma samples must be diluted at least 15-fold in diluent.
  - The optimal sample dilution should be determined empirically. We suggest an initial sample dilution of 50-fold.
  - Optimum results are achieved if, at each step, reagents are pipetted into the wells of the microtiter plate within 5 minutes.

- Limitations of the procedure
  - Reliable and reproducible results will be obtained when the assay procedure is carried out with a complete understanding of and in accordance with the instructions detailed above.
  - The wash procedure is critical. Insufficient washing will result in poor precision and falsely elevated absorbance readings.
Assay Protocol

Reagent Preparation

- Wash Solution Preparation
  The wash solution is provided as a 20x stock. Prior to use dilute the contents of the bottle (50 mL) with 950 mL of distilled or deionized water.

- Standard Preparation
  1. Reconstitute the lyophilized mouse anti-SRBC IgG standard stock with diluent as described on the standard vial label (the reconstituted standard should be aliquoted and frozen at -20°C after reconstitution if additional use is intended).
  2. Label 6 polypropylene or glass tubes as 100, 50, 25, 12.5, 6.25, and 3.125 u/mL.
  3. In the tube labeled 100 u/mL, prepare a 100 u/mL stock by mixing the volume of reconstituted standard stock with the volume of diluent detailed on the reference standard stock vial label.
  4. Dispense 250 μL of diluent into the tubes labeled 50, 25, 12.5, 6.25, and 3.125 u/mL.
  5. Prepare a 50 u/mL standard by diluting and mixing 250 μL of the 100 u/mL standard with 250 μL of diluent in the tube labeled 50 u/mL.
  6. Similarly prepare the 25, 12.5, 6.25, and 3.125 u/mL standards by serial dilution.

Sample Preparation

Anti-SRBC IgG is present in mouse serum or plasma from SRBC immunized animals at concentrations in excess of 500 u/mL. In order to obtain values within the range of the standard curve, we suggest that samples initially be diluted 50 fold using the following procedure:
  1. For each test sample dispense 294 μL of diluent into separate tubes.
  2. Pipette and mix 6 μL of the serum/plasma sample into a tube containing 294 μL of diluent. This provides a 50 fold diluted sample.
  3. Repeat this procedure for each sample to be tested Important. Do not use dilutions lower than 15 fold.

Assay Procedure

1. Secure the desired number of coated wells in the holder.
2. Dispense 100 μL of standards and diluted samples into the wells (we recommend that samples be tested in duplicate).
3. Incubate on an orbital micro-plate shaker at 100-150 rpm at room temperature (18-25°C) for 45 minutes.
4. Aspirate the contents of the microtiter wells and wash the wells 5 times with 1x wash solution using a plate washer (400 μL/well). The entire wash procedure should be performed as quickly as possible.
5. Strike the wells sharply onto absorbent paper or paper towels to remove all residual wash buffer.
6. Add 100 μL of enzyme conjugate reagent into each well.
7. Incubate on an orbital micro-plate shaker at 100-150 rpm at room temperature (18-25°C) for 45 minutes.
8. Wash as detailed in 4 to 5 above.
9. Dispense 100 μL of TMB Reagent into each well.
10. Gently mix on an orbital micro-plate shaker at 100-150 rpm at room temperature (18-25°C) for 20 minutes.
11. Stop the reaction by adding 100 μL of Stop Solution to each well.
12. Gently mix. It is important to make sure that all the blue color changes to yellow.
13. Read the optical density at 450 nm with a microtiter plate reader within 5 minutes.
Data Analysis

Calculation of Results

1. Calculate the average absorbance value ($A_{450}$) for each set of reference standards and samples.
2. Construct a standard curve by plotting the mean absorbance obtained from each reference standard against its concentration in $\mu$/mL on linear graph paper, with absorbance values on the vertical or Y-axis and concentrations on the horizontal or X-axis.
3. Using the mean absorbance value for each sample, determine the corresponding concentration of anti-SRBC IgG in $\mu$/mL from the standard curve.
4. Multiply the derived concentrations by the dilution factor to determine the actual concentration of anti-SRBC IgG in the serum/plasma sample.
5. PC graphing software may be used for the above steps.
6. If the $OD_{450}$ values of samples fall outside the standard curve when tested at a dilution of 50, samples should be diluted appropriately and re-tested (do not use dilutions lower than 15 fold).

A typical standard curve with optical density readings at 450 nm on the Y axis against anti-SRBC IgG concentrations on the X axis is shown below. This curve is for the purpose of illustration only and should not be used to calculate unknowns. Each user should obtain his or her data and standard curve in each experiment.

<table>
<thead>
<tr>
<th>Anti-SRBC IgG (u/mL)</th>
<th>Absorbance (450 nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>100</td>
<td>2.556</td>
</tr>
<tr>
<td>50</td>
<td>1.423</td>
</tr>
<tr>
<td>25</td>
<td>0.782</td>
</tr>
<tr>
<td>12.5</td>
<td>0.461</td>
</tr>
<tr>
<td>6.25</td>
<td>0.292</td>
</tr>
<tr>
<td>3.125</td>
<td>0.201</td>
</tr>
</tbody>
</table>
Resources

Reference

Plate Layout