KAMIYA BIOMEDICAL COMPANY

Salmon and Trout IgM ELISA

For the quantitative determination of IgM in salmon and trout serum

Cat. No. KT-1913

For Research Use Only.

PRODUCT INFORMATION

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PRODUCT

The ***K-ASSAY®*** Salmon and Trout IgM ELISA is an enzyme immunoassay for the quantitative determination of IgM in salmon and trout serum. For research use only.

INTRODUCTION

IgM is the most abundant immunoglobulin in trout and salmon serum. It is a tetramer; each subunit is comprised of two ~75 kDa heavy chains and two ~25 kDa light chains. In trout, levels range from 2 – 10 mg/mL, levels in salmon are approximately 1 mg/mL. It has been reported that total IgM levels increase in salmon during infection.

PRINCIPLE

The assay uses a monoclonal IgM antibody that recognizes the heavy chain of rainbow trout and Atlantic salmon IgM. The unconjugated antibody is coated on wells of a microtiter plate and used for capture. A horseradish peroxidase (HRP) conjugate is used for detection. Calibrators and diluted samples (100 µL) are incubated in the antibody coated microtiter wells for 45 minutes. After washing the wells, HRP-conjugate (100 µL) is added and incubated for 45 minutes. If IgM molecules are present, they are sandwiched between the capture and detection antibodies. The wells are then washed to remove unbound HRP-conjugate. TMB is added and incubated for 20 minutes. If IgM is present, a blue color develops. Color development is stopped by addition of Stop Solution, changing the color to yellow. Absorbance is measured at 450 nm. The concentration of IgM is proportional to absorbance and is derived from a calibration curve.

COMPONENTS

- Anti-IgM coated plate (12 x 8-well strips)
- HRP conjugate, 11 mL
- IgM stock calibrator, 3 vials
- 20x Wash Solution, 50 mL
- Diluent, 2 x 50 mL
- TMB, 11 mL
- Stop Solution, 11 mL

MATERIALS REQUIRED BUT NOT PROVIDED

- Pipettors and tips
- Distilled or de-ionized water
- Polypropylene tubes or 96-well polystyrene plates
- Vortex mixer
- Absorbent paper or paper towels
- Plate incubator/shaker
- Plate washer
- Plate reader capable of measuring absorbance at 450 nm
- Graphing software

GENERAL INSTRUCTIONS

1. All reagents should be allowed to reach room temperature before use.
2. Reliable and reproducible results will be obtained when the assay is conducted with a complete understanding of the instructions and with adherence to good laboratory practice.
3. It is important that calibrators and samples be added to the ELISA plate quickly. If testing large numbers of samples, rather than pipetting calibrators and samples from individual tubes into the ELISA plate, we recommend the following. First, pipette an excess volume of calibrators and samples into wells of a blank polystyrene 96-well plate. Then use an 8 or 12-channel multi-pipettor to quickly transfer 100 µL aliquots to the appropriate wells of the antibody-coated plate.
4. The wash procedure is critical. Insufficient washing will result in poor precision and falsely elevated absorbance readings.
5. Laboratory temperature will influence absorbance readings. The assay was calibrated using a shaking incubator set at 150 rpm and 25℃. Performing the assay at lower temperatures and mixing speeds may result in lower absorbance values.

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 de-ionized water. Unused wash buffer may be stored at 4°C for one week.

DILUENT PREPARATION

The diluent is formulated for measurement of IgM in salmon and trout serum. It is supplied ready to use. DO NOT substitute other buffers.

CALIBRATOR

1. The stock is lyophilized. Reconstitute it with the volume of diluent shown on the vial label and prepare the 250 ng/mL calibrator as described.
2. Label seven polypropylene tubes as 125, 62.5, 31.25, 15.63, 7.81, 3.91, and 1.95 ng/mL. Dispense 0.25 mL of diluent into each.
3. Pipette 0.25 mL of the 250 ng/mL IgM calibrator into the tube labeled 125 ng/mL and mix. This provides the 125 ng/mL IgM calibrator.
4. Similarly prepare the remaining calibrators by two-fold serial dilution.

Unused reconstituted IgM is stable overnight in the refrigerator.

SAMPLE PREPARATION

In studies, we found IgM levels ranging from 1 to 6 mg/mL in rainbow trout serum. In Atlantic salmon, we found that levels ranged from 0.1 to 0.3 mg/mL. We suggest that trout serum be evaluated at a dilution of 40,000-fold and salmon serum at a dilution of 2,000-fold. Suggested dilution strategies are listed below. Ideally, dilutions should be performed in polystyrene 96-well plates (not provided). This allows quick and easy transfer of diluted samples to the antibody-coated plate using 8- or 12-channel multi-pipettors.

40,000-fold Dilution Strategy

1. Pipette 198 µL into two wells and 187.5 µL into a third well.
2. Pipette 2.0 µL of serum into the first well containing 198 µL of diluent and mix. This provides a 100-fold dilution.
3. Pipette 2.0 µL of the 100-fold diluted sample into the second well containing 198 µL of diluent and mix. This provides a 10,000-fold dilution.
4. Pipette 62.5 µL of the 10,000-fold diluted sample into the third well containing 187.5 µL of diluent and mix. This provides a 40,000-fold dilution.

2,000-fold Dilution Strategy

1. Pipette 198 µL into one well and 237.5 µL into a second well.
2. Pipette 2.0 µL of serum into the first well containing 198 µL of diluent and mix. This provides a 100-fold dilution.
3. Pipette 12.5 µL of the 100-fold diluted sample into the second well containing 237.5 µL of diluent and mix. This provides a 2,000-fold dilution.

ASSAY PROCEDURE

1. Secure the desired number of 8-well strips in the cassette. Unused strips should be stored in a sealed bag with desiccant at 4°C.
2. Dispense 100 µL of calibrators and samples into the wells.
3. Incubate on a plate shaker at 150 rpm and 25°C for 45-minutes.
4. Empty and wash the microtiter wells 5x with 1x wash solution using a plate washer (400 µL/well).
5. Dispense 100 µL of HRP conjugate into the wells.
6. Incubate on a plate shaker at 150 rpm and 25°C for 45-minutes.
7. Strike the wells sharply onto absorbent paper or paper towels to remove all residual droplets.
8. Dispense 100 µL of TMB into each well.
9. Incubate on an orbital micro-plate shaker at 150 rpm at 25°C for 20 minutes.
10. After 20-minutes, stop the reaction by adding 100 µL of Stop solution to each well.
11. Gently mix. It is important to make sure that all the blue color changes to yellow.
12. Read absorbance at 450 nm with a plate reader within 5 minutes.

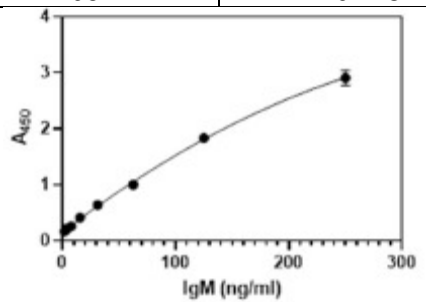
CALCULATION OF RESULTS

1. Using curve fitting software, construct a calibration curve by plotting absorbance values of the calibrators versus the IgM concentration.
2. Fit the calibration curve using graphing software. We suggest using a second order polynomial (quadratic) equation.
3. Derive the concentration by the dilution factor to determine the concentration in the sample.
4. If the absorbance values of samples fall outside the calibration curve, samples should be diluted appropriately and re-tested.

TYPICAL CALIBRATION CURVE

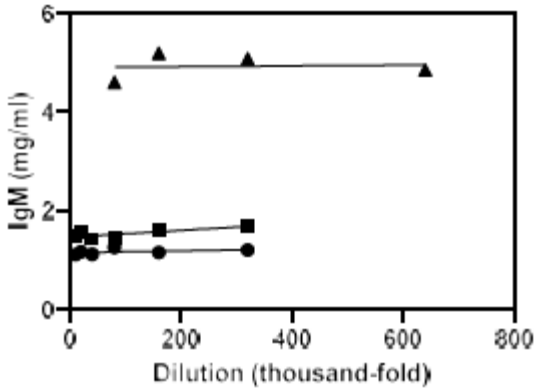
A typical calibration curve is shown below. This curve is for illustration only.

IgM (ng/mL)	A_{450}
250	2.909
125	1.835
62.5	1.003
31.25	0.641
15.63	0.416
7.81	0.263
3.91	0.221
1.95	0.173



ASSAY PERFORMANCE

Linearity: To assess the linearity of the assay, three rainbow trout serum samples with IgM concentrations of 1.17, 1.54 and 4.77 mg/mL were serially diluted to produce values within the dynamic range of the assay.



STORAGE

The kit should be stored at 4℃ and the microtiter plate should be kept in a sealed bag with desiccant. The kit will remain stable until the expiration date.

FOR RESEARCH USE ONLY

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