

KAMIYA BIOMEDICAL COMPANY

Rat CK-MM ELISA

For the quantitative determination of creatine kinase MM (CK-MM) in rat serum or plasma.

Cat. No. KT-551

For Research Use Only.

PRODUCT INFORMATION

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PRODUCT

The **K-ASSAY®** Rat CK-MM ELISA is an enzyme immunoassay for the quantitative determination of CK-MM in rat serum or plasma. For research use only.

INTRODUCTION

Creatine kinase (CK) is a key metabolic enzyme. It is a dimer of two subunits, each with molecular weights of approximately 43 kDa. Two different subunits occur, M and B. The CK holoenzyme therefore exists as MM and BB homodimers and an MB heterodimer. The MM and BB isoforms are expressed primarily in skeletal muscle and brain respectively while both MB and MM are expressed in heart. The relatively high expression levels of the MB isoform in heart explain its use as an established biomarker for heart disease. Likewise, CK-MM can be used as a specific biomarker for skeletal muscle injury.

PRINCIPLE

The **K-ASSAY®** Rat CK-MM ELISA is a solid phase enzyme-linked immunosorbent assay (ELISA). The assay uses affinity purified CK-MM antibodies for solid phase immobilization and a horseradish peroxidase (HRP) conjugated CK-MM monoclonal antibody for detection. Calibrators and diluted test samples are incubated in the microtiter wells for 45 minutes. The microtiter wells are subsequently washed and HRP conjugate is added and incubated for 45 minutes. As a result, CK-MM molecules are sandwiched between the immobilization and detection antibodies. 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 CK-MM is proportional to the optical density of the test sample and actual concentrations are determined by reference to a calibration curve.

COMPONENTS

- Anti-rat CK-MM antibody coated microtiter plate with 96 wells (provided as 12 detachable strips of 8)
- 2X HRP Conjugate, 6 mL
- Reference rat CK-MM stock (lyophilized)
- Wash Buffer (20X), 50 mL
- Diluent, 50 mL
- TMB Reagent (One-Step), 11 mL
- Stop Solution (1N HCl), 11 mL

MATERIALS REQUIRED BUT NOT PROVIDED

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

GENERAL INSTRUCTIONS

All reagents should be allowed to reach room temperature (18-25°C) before use.

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.

HRP CONJUGATE PREPARATION

The HRP conjugate is provided as a 2X stock. Prior to use estimate the final volume of HRP conjugate required for your assay and dilute one (1) volume of the 2X stock with one (1) volume of diluent. Typically, we prepare 1 mL of conjugate for each 8-well strip used in the assay. This is prepared by mixing 0.5 mL of 2X HRP conjugate with 0.5 mL of diluent. Mix gently prior to use. Prepare the working conjugate no more than one hour in advance.

CALIBRATOR PREPARATION

1. The reference CK-MM stock is provided in lyophilized form. Reconstitute with diluent as directed on the vial label and mix gently until dissolved. **(the reconstituted calibrator remains stable for 1 day at 4 °C).**
2. Label 5 polypropylene or glass tubes as 50, 25, 12.5, 6.25 and 3.13 ng/mL.
3. Prepare the 50 ng/mL calibrator as described on the reference stock vial label.
4. Dispense 250 μ L of diluent into the tubes labeled 25, 12.5, 6.25 and 3.13 ng/mL.
5. Pipette 250 μ L of the 50 ng/mL CK-MM calibrator into the tube labeled 25 ng/mL and mix. This provides the working 25 ng/mL CK-MM calibrator.
6. Similarly prepare the 12.5, 6.25 and 3.13 ng/mL calibrators by serial dilution.

SAMPLE PREPARATION

General Note: CK-MM is usually below the detection level of the kit in samples from animals without skeletal muscle injury. However, after skeletal muscle injury levels can increase to 8,000 ng/mL or greater. Undiluted serum or plasma must not be used in the ELISA. We recommend that samples be diluted at least 10-fold or more. This eliminates interference caused by serum/plasma components. Assuming that a dilution of 10-fold is used; samples should be treated as follows:

1. Dispense 225 μ L of diluent into separate tubes.
2. Pipette and mix 25 μ L of the serum/plasma sample into the tube containing 225 μ L of diluent. This provides a 10 fold diluted sample.
3. Repeat this procedure for each sample to be tested.

ASSAY PROCEDURE

1. Secure the desired number of coated wells in the holder.
2. Dispense 100 μ L of calibrators 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. Remove the incubation mixture using either a plate washer or by flicking plate contents into an appropriate Bio-waste container.
5. Wash and empty the microtiter wells 5 times with 1X wash solution. This should preferentially be performed using a plate washer (400 μ L/well). If a plate washer is not available use a squirt bottle. The entire wash procedure should be performed as quickly as possible.
6. Strike the wells sharply onto adsorbent paper or paper towels to remove all residual droplets.
7. Add 100 μ L of enzyme conjugate reagent into each well.
8. Incubate on an orbital micro-plate shaker at 100-150 rpm at room temperature (18-25°C) for 45 minutes.
9. Wash as detailed in 4 to 5 above.
10. Strike the wells sharply onto absorbent paper or paper towels to remove residual droplets.
11. Dispense 100 μ L of TMB Reagent into each well.
12. Gently mix on an orbital micro-plate shaker at 100-150 rpm at room temperature (18-25°C) for 20 minutes.
13. Stop the reaction by adding 100 μ L of Stop Solution to each well.
14. Gently mix. *It is important to make sure that all the blue color changes to yellow.*
15. Read the optical density at 450 nm with a microtiter plate reader within 5 minutes.

CALCULATION OF RESULTS

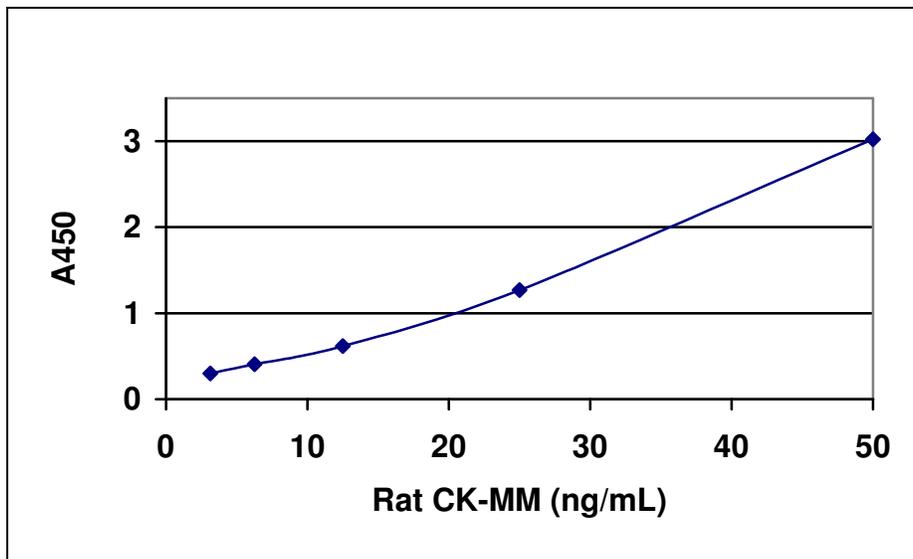
1. Calculate the average absorbance values (A_{450}) for each set of reference calibrators, and samples.
2. Construct a calibration curve by plotting the mean absorbance obtained from each reference calibrator against its concentration in ng/mL on linear graph paper, with absorbance values on the vertical 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 CK-MM in ng/mL from the calibration curve.
4. Multiply the derived concentrations by the dilution factor to determine the actual concentration of CK-MM in the serum/plasma sample.
5. PC graphing software may be used for the above steps. We recommend a point-to-point "fit" of the data because of the concave shape of the calibration curve. Alternatively, fitting to a polynomial second order equation may work well.
6. If the A_{450} values of samples exceed that of the high calibrator the samples should be further diluted and re-tested.

Samples with absorbance values below the lowest calibrator should be assigned a value of $\leq 3.13 \times A$ ng/mL, where A represents the dilution factor used for the sample.

TYPICAL CALIBRATION CURVE

A typical calibration curve with optical density readings at 450 nm on the Y axis against CK-MM 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 calibration curve in each experiment.

CK-MM (ng/mL)	Absorbance (450 nm)
50	3.021
25	1.269
12.5	0.618
6.25	0.406
3.13	0.299



STORAGE

The kit should be stored at 4°C and the microtiter strips should be kept in a sealed bag with desiccant to minimize exposure to damp air. Test kits will remain stable until the expiration date provided that the components are stored as described above.

LIMITATIONS OF THE PROCEDURE

1. Do not use samples at a dilution less than 10-fold (i.e., do not use a dilution of 5-fold). High serum content of the samples can interfere with the ELISA.
2. 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.
3. The wash procedure is critical. Insufficient washing will result in poor precision and falsely elevated absorbance readings.

SPECIFICITY

Rat CK-BB and rat CK-MB were not available when this kit was developed and specificity was therefore investigated using purified human CK-MM, CK-MB and CK-BB. Human CK-MM was strongly recognized but CK-BB was not recognized at concentrations up to 10 micrograms/mL. The ELISA was ~400-fold more selective for CK-MM compared to CK-MB

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