

RAT C-REACTIVE PROTEIN (CRP) ELISA

Life Diagnostics, Inc., Catalog Number: CRP-2

INTRODUCTION

CRP is an acute phase protein that is elevated in serum due to injury, infection, or disease. The level of induction varies considerably with species, ranging from approximately two-fold in mice and rats to 1000-fold or more in dogs and humans.

PRINCIPLE OF THE ASSAY

The assay uses affinity purified rat CRP antibodies for solid phase (microtiter wells) immobilization and horseradish peroxidase (HRP) conjugated rat CRP antibodies for detection. Standards and diluted samples are incubated in the microtiter wells for 45 minutes. The wells are subsequently washed. HRP conjugate is added and incubated for 45 minutes. This results in CRP molecules being sandwiched between the immobilization and detection antibodies. The wells are then washed to remove unbound HRP-conjugate and TMB is added and incubated for 20 minutes. If CRP is present a blue color develops. Color development is stopped by the addition of Stop solution, changing the color to yellow. Absorbance is measured at 450 nm. The concentration of CRP is proportional to absorbance and is derived from a standard curve.

MATERIALS AND COMPONENTS

Materials provided with the kit:

- CRP antibody coated 96-well plate (12 x 8-well strips)
- HRP Conjugate, 11 ml
- CRP stock (lyophilized)
- 20x Wash solution: TBS50-20, 50 ml
- 10x Diluent: YD25-10, 25 ml
- TMB: TMB11-1, 11 ml
- Stop solution: SS11-1, 11 ml

Materials required but not provided:

- Pipettors and tips
- Distilled or deionized water
- Polypropylene or glass tubes
- Vortex mixer
- Absorbent paper or paper towels
- Plate incubator/shaker
- Plate washer
- Plate reader capable of measuring absorbance at 450 nm.
- Curve fitting software

STORAGE

The unused kit should be stored at 4°C and the microtiter plate should be kept in a sealed bag with desiccant. Kits will remain stable for six months from the date of purchase.

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 carried out with a complete understanding of the instructions and with adherence to good laboratory practice.
3. The wash procedure is critical. Insufficient washing will result in poor precision and falsely elevated absorbance readings.
4. Laboratory temperature will influence absorbance readings. Our EISA kits are calibrated using shaking incubators set at 150 rpm and 25°C. Performance of the assay at lower temperatures will result in lower absorbance values.

DILUENT PREPARATION

The diluent is provided as a 10x stock. Prior to use estimate the final volume of diluent required for your assay and dilute one volume of the 10x stock with nine volumes of distilled or deionized water.

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. The CRP stock is provided lyophilized. Add the volume of distilled or de-ionized water indicated on the vial label and mix gently until dissolved (*the reconstituted standard remains stable for at least 10 days at 2-8°C but should be aliquoted and frozen at -20°C after reconstitution if use beyond this time is intended*).
2. Label 5 polypropylene or glass tubes as 10, 5, 2.5, 1.25 and 0.625 ng/ml.
3. In the tube labeled 10 ng/ml prepare the 10 ng/ml standard as detailed on the stock vial label.
4. Dispense 250 μ l of diluent into the tubes labeled 5, 2.5, 1.25, and 0.625 ng/ml.
5. Prepare the 5 ng/ml standard by mixing 250 μ l of the 10 ng/ml standard with 250 μ l of diluent in the tube labeled 5 ng/ml.
6. Similarly prepare the remaining standards by two-fold serial dilution.

SAMPLE PREPARATION

CRP is present in normal rat serum at a concentration of approximately 0.5 mg/ml. To obtain values within the range of the standard curve we suggest that samples be diluted 100,000-fold using the following procedure for each sample to be tested:

1. Dispense 998 μ l and 497.5 μ l of 1x diluent into separate tubes.
2. Pipette and mix 2 μ l of the serum/plasma sample into the tube containing 998 μ l of diluent. This provides a 500-fold diluted sample.
3. Mix 2.5 μ l of the 500-fold diluted sample with the 497.5 μ l of diluent in the second tube. This provides a 100,000-fold dilution of the sample.

- Repeat this procedure for each sample to be tested.

ASSAY PROCEDURE

- Secure the desired number of 8-well strips in the holder. Unused strips should be stored in the re-sealed bag with desiccant at 4°C for future use.
- Dispense 100 µl of standards and samples into the wells (we recommend that standards and samples be run in duplicate).
- Incubate on an orbital micro-plate shaker at 150 rpm and 25°C for 45 minutes.
- Empty and wash the microtiter wells 5x with 1x wash solution using a plate washer (400 µl/well).
- Strike the wells sharply onto absorbent paper or paper towels to remove all residual droplets.
- Add 100 µl of HRP-conjugate into each well.
- Incubate on a plate shaker at 150 rpm and 25°C for 45 minutes.
- Wash as detailed above.
- Strike the wells sharply onto absorbent paper or paper towels to remove residual droplets.
- Dispense 100 µl of TMB into each well.
- Incubate on an orbital micro-plate shaker at 150 rpm at 25°C for 20 minutes.
- After 20-minutes, stop the reaction by adding 100 µl of Stop solution to each well.
- Gently mix. It is important to make sure that all the blue color changes to yellow.
- Read absorbance at 450 nm with a plate reader within 5 minutes.

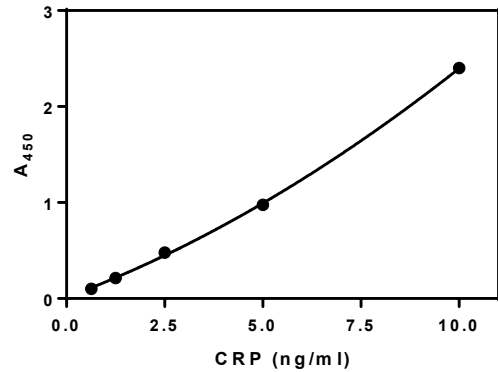
CALCULATION OF RESULTS

- Using curve fitting software, construct a standard curve by plotting absorbance values of the standards versus concentration.
- Fit the standard curve to an appropriate model and derive the concentration of the samples (we recommend using a single site, total and nonspecific binding model).
- Multiply the derived concentration by the dilution factor to determine the actual concentration in the serum sample.
- If the A_{450} values of samples fall outside the standard curve, samples should be diluted appropriately and re-tested.

TYPICAL STANDARD CURVE

A typical standard curve is shown below. This curve is for illustration only and should not be used to calculate unknowns.

CRP (ng/ml)	Absorbance (450 nm)
10	2.4
5	0.975
2.5	0.479
1.25	0.213
0.625	0.102



Rev 12082020

For technical assistance please email us at techsupport@lifediagnosics.com