

Camel Total Antioxidant Capacity (T-AOC) ELISA Kit

to determine Camel Total Antioxidant Capacity ELISA Kit in Serum, Blood Plasma, Saliva, Urine, Tissue Liquid Samples or related Biological Solutions.

INSTRUCTION MANUAL

FOR ELISA KIT No: LT879910IBKCCBA



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INTENDED USE

This kit is used to determine the Camel Total Antioxidant Capacity (T-AOC) ELISA Kit in the sample of serum, blood plasma, saliva, urine, tissue liquid samples or related biological solutions. For *In vitro* use only.

MANUAL VERSION 1.0IBF

INTRODUCTION

ASSAY PRINCIPLE

Camel T-AOC ELISA Kit employs a sandwich ELISA to quantitate Camel T-AOC, in samples.

Add Standard and Sample to the wells that pre-coated with antibody, then add HRP-Conjugate reagent to form an immune complex, incubation, by incubation and washing, removal of unbound enzyme, and then add the substrate A and B, then the solution will turn blue and finally change into yellow with the effect of acid. The color absorbance is positively correlated with the concentration of Total Antioxidant Capacity (T-AOC).

CHARACTERISTICS

- This Kit allows for the determination of Camel T-AOC concentrations in Camel serum, cell culture supernates and other biological fluids.
- Detection range: : 1U/mL -24U/mL
- The minimum detectable dose (MDD) of Camel T-AOC is typically less than 0.1U/MI
- Four samples of known concentration were tested twenty times on one plate to assess intra-assay precision. The CV (%) <10%.
- Three samples of known concentration were tested in twenty separate assays to assess inter-assay precision. Assays were performed by at least three technicians using two lots of components. The CV (%) <15%.
- To assess linearity of the assay, samples containing and/or spiked with high concentrations of Camel T-AOC were diluted with the appropriate calibrator diluent to produce samples with values within the dynamic range of the

assay. Linear regression analysis of samples versus the expected concentration yielded a correlation coefficient of 0.99.

- Camel T-AOC Kit has high sensitivity and excellent specificity for detection of Camel T-AOC. No significant cross-reactivity or interference was observed.

PRODUCT INFORMATION

MATERIALS SUPPLIED AND STORAGE CONDITIONS

Store kit reagents at 2-8 °C for 6 months.

Components	96 T	Storage conditions
Microelisa stripplate	96 wells	2-8 °C ¹
HRP-Conjugated detection antibody	10 ml	2-8 °C
Sample diluent	6 ml	2-8 °C
Standard A	0U/mL	2-8 °C
Standard B	1.5U/mL	2-8 °C
Standard C	3U/mL	2-8 °C
Standard D	6U/mL	2-8 °C
Standard E	12U/mL	2-8 °C
Standard F	24U/mL	2-8 °C
Chromogen solution A	6 ml	2-8 °C
Chromogen solution B	6 ml	2-8 °C
Stop solution	6 ml	2-8 °C
Wash buffer, 20X	25 ml	2-8 °C
Plate covers	2	RT
Booklet	1	RT
Sealed bags	1	RT

¹Return unused wells to the foil pouch containing the desiccant pack. Reseal along entire edge of the zip-seal. May be stored for up to 1 month at 2-8 °C.

MATERIALS REQUIRED BUT NOT SUPPLIED

1. 37 °C incubator
2. Microplate reader
3. Precision pipettes and tips
4. Distilled water
5. Disposable tubes for sample dilution
6. Absorbent paper

ASSAY PROTOCOL

SAMPLE COLLECTION AND STORAGE

The sample collection and storage conditions listed below are intended as general guidelines. Sample stability has not been evaluated.

Tissue homogenates - For general information, hemolysis blood may affect the result, so you should rinse the tissues with ice-cold PBS (0.01M, pH=7.4) to remove excess blood thoroughly. Tissue pieces should be weighed and then minced to small pieces which will be homogenized in PBS with a glass homogenizer on ice. (The volume depends on the weight of the tissue, 9mL PBS would be appropriate to 1 gram tissue pieces. Some protease inhibitor is recommended to add into the PBS.) To further break the cells, you can sonicate the suspension with an ultrasonic cell disrupter or subject it to freeze-thaw cycles. The homogenates are then centrifuged for 20 minutes at 2000-3000 RPM to collect the supernate.

Cell culture supernatants and other biological fluids - Centrifuge samples for 20 minutes at 2000-3000 RPM. Remove particulates and assay immediately or store samples in aliquot at -20 °C or -80 °C for later use. Avoid repeated freeze/thaw cycles.

Serum - Use a serum separator tube and allow samples to clot for two hours at room temperature or overnight at 4 °C before centrifugation for 20 minutes at 2000-3000 RPM. Assay freshly prepared serum immediately or store samples in aliquot at -20 °C or -80 °C for later use. Avoid repeated freeze/thaw cycles.

Plasma - Collect plasma using EDTA or heparin as an anticoagulant. Centrifuge

samples for 20 minutes at 2000-3000 RPM at 2-8 °C within 30 minutes of collection. Remove plasma and assay immediately or store samples in aliquot at -20 °C or -80 °C for later use. Avoid repeated freeze/thaw cycles.

Urine: Collect urine in a clean sterile tube. Centrifuge at 2000-3000 RPM for approximately 20 minutes. Collect the supernatant carefully. If sedimentation occurs during storage, centrifugation should be performed again.

Note: Samples should be centrifuged adequately and no haemolysis or granule was allowed.

REAGENT PREPARATION

Bring all reagents to room temperature before use. If crystals have formed in the Buffer Concentrates, warm them gently until they completely dissolved.

Wash buffer - Dilute with Distilled or deionized water 1:20.

Standard – Standards are ready to use.

ASSAY PROCEDURE

1. Prepare all reagents before starting assay procedure. It is recommended that all Standards and Samples be added in duplicate to the microplate.
Note: If samples generate values higher than the highest standard, please dilute the samples with Sample Diluent and repeat the assay.
2. Add standard: Set Standard wells, testing sample wells. Add standard 50µl to standard well.
3. Add Sample: Add sample diluent 40 µl to testing sample well. Then add sample 10µl to testing sample well; for Blank well don't add anything.
4. Add 100 µl HRP-Conjugate to each well, except blank well. Cover with a plate cover and incubate for 60 minutes at 37 °C.

5. Aspirate each well and wash, repeating the process four times for a total of five washes with soak time of 1 min. Wash by filling each well with Wash buffer (350 µl) using a squirt bottle, manifold dispenser or auto washer by aspirating or decanting. Invert the plate and blot it against clean paper towels.

6. Automatic washing: Each well inject into the wash solution 350µL, soak 1min, wash plate 5 times.)

7. Add 50 µl chromogen solution A and 50 µl chromogen solution B to each well. Gently mix and incubate for 15 minutes at 37°C. Protect from light.

8. Add 50µl Stop Solution to each well. The color in the wells should change

from blue to yellow. If the color in the wells is green or the color change does not appear uniform, gently tap the plate to ensure thorough mixing.

9. Read the Optical Density (O.D.) at 450 nm using a microtiter plate reader within 15 minutes.

Summary of operating procedures

Prepare reagents, samples and standards



Add prepared samples and standards, and HRP, 37°C incubation for 60 minutes



Wash the plate 5 times, add chromogen solution A, B, 37°C developing color for 15 minutes



Add the stop solution

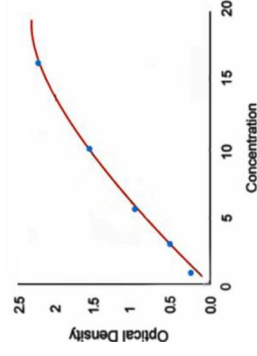


Read the OD value within 15 minutes

CALCULATION

Average the duplicate readings for each standard, control, and sample and subtract the average zero standard optical density (O.D.). Create a standard curve by reducing the data using computer software capable of generating a four parameter logistic (4-PL) curve fit. As an alternative, construct a standard curve by plotting

the mean absorbance for each standard on the y-axis against the concentration for each standard on the x-axis and draw a best fit curve through the points on the graph. The data may be linearized by plotting the log of the Camel T-AOC concentration versus the log of the O.D. and the best fit line can be determined by regression analysis. This procedure will produce an adequate but less precise fit of the data. If samples have been diluted, the concentration read from the standard curve must be multiplied by the dilution factor (for e.g. it is 5 in this case).



PRECAUTIONS

1. Do not mix or substitute reagents with those from other lots or sources.
2. This assay is designed to eliminate interference by other factors present in biological samples. Until all factors have been tested in the ELISA Immunoassay, the possibility of interference cannot be excluded.
3. Any variation in standard diluent, operator, pipetting technique, washing technique, incubation time or temperature, and kit age can cause variation in binding.
4. When mixing or reconstituting protein solutions, always avoid foaming.
5. To avoid cross-contamination, change pipette tips between additions of each standard level, between sample additions, and between reagent additions. Also, use

separate reservoirs for each reagent.

6. To ensure accurate results, proper adhesion of plate sealers during incubation steps is necessary.
7. When using an automated plate washer, adding a 30 second soak period following the addition of Wash Buffer, and/or rotating the plate 180 degrees between wash steps may improve assay precision.
8. Chromogen Solution is easily contaminated. If bluish prior to use, do not use.
9. Stop Solution should be added to the plate in the same order as the Chromogen solution. The color developed in the wells will turn from blue to yellow upon addition of the Stop Solution. Wells that are green in color indicate that the Stop Solution has not mixed thoroughly with the Chromogen solution.
10. Serum and plasma should be handled as potentially hazardous and capable of transmitting disease. Disposable gloves must be worn during the assay procedure, since no known test method can offer complete assurance that products derived from Rat blood will not transmit infectious agents. Therefore, all blood derivatives should be considered potentially infectious and good laboratory practices should be followed.
11. All samples should be disposed of in a manner that will inactivate viruses.
12. Liquid Waste: Add sodium hypochlorite to a final concentration of 1.0%. The waste should be allowed to stand for a minimum of 30 minutes to inactivate the viruses before disposal.

VALIDITY & STORAGE: As per CoA.