



## **Ochratoxin A (OTA) ELISA Kit**

**Catalog Number. CSB-EFD027449**

**This immunoassay kit allows for the in vitro quantitative/qualitative determination of Ochratoxin A concentrations in feed, grain, cereal.**

This package insert must be read in its entirety before using this product.

### **If You Have Problems**

### **Technical Service Contact information**

Phone: 86-27-87582341

Fax: 86-27-87196150

Email: [tech@cusabio.com](mailto:tech@cusabio.com)

Web: [www.cusabio.com](http://www.cusabio.com)

In order to obtain higher efficiency service, please ready to supply the lot number of the kit to us (found on the outside of the box).

## **INTRODUCTION**

Ochratoxin A (OTA) is a toxic metabolite produced by *Aspergillus flavus* and *Penicillium* genera. Ochratoxin A has a strong liver and kidney toxicity, and has teratogenicity, mutagenicity and carcinogenicity. Ochratoxin A was considered to be the second important and harmful mycotoxin in the family following with Aflatoxin flavus. As the Ochratoxin A producing fungi exist widely in nature, especially higher yield in cereals and cereals products, coffee, cocoa contaminated with Ochratoxin A, it has been potential threat for human health and development of animal husbandry.

This kit is a detection product developed based on ELISA technology, with operation time as short as 50 min and a sensitivity of 0.15 µg/kg, and linear range from 0.15 µg/kg to 4.05 µg/kg.

## **PRINCIPLE OF THE ASSAY**

This assay employs the competitive inhibition enzyme immunoassay technique. The microtiter plate provided in this kit has been pre-coated with Ochratoxin A toxin antigen. Standards or samples are added to the appropriate microtiter plate wells with a Horseradish Peroxidase (HRP) and antibody of Ochratoxin A. The competitive inhibition reaction is launched between pre-coated Ochratoxin A and Ochratoxin A in standards or samples with the Ochratoxin A special antibody. A substrate solution is added to the wells and the color develops in opposite to the amount of Ochratoxin A in the standards or samples. The color development is stopped and the intensity of the color is measured.

## **DETECTION RANGE**

0.15µg/kg~4.05µg/kg

## **SENSITIVITY**

The minimum detectable dose of the kit is typically less than 0.15 µg/kg.

The sensitivity of this assay, or Lower Limit of Detection (LOD) was defined as the lowest concentration that could be differentiated from zero.

### **CROSS-REACTION RATE**

Ochratoxin A	100%
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### **RECOVERY RATE**

Feed/Grain/Cereal	70%-100%
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### **LIMIT OF DETECTION**

Feed/Grain/Cereal	2 µg/kg
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### **PRECISION**

#### **Intra-assay Precision (Precision within an assay): CV%<10%**

Three samples of known concentration were tested twenty times on one plate to assess.

#### **Inter-assay Precision (Precision between assays): CV%<10%**

Three samples of known concentration were tested in twenty assays to assess.

### **LIMITATIONS OF THE PROCEDURE**

- The kit should not be used beyond the expiration date on the kit label.
- Do not mix or substitute reagents with those from other lots or sources.
- Any variation in operator, pipetting technique, washing technique, incubation time or temperature, and kit age can cause variation in binding.
- This assay is designed to eliminate interference by soluble receptors, binding proteins, and other factors present in biological samples. Until all factors have been tested in the Immunoassay, the possibility of interference cannot be excluded.

## **MATERIALS PROVIDED**

<b>Reagent</b>	<b>Quantity(96T)</b>	<b>Quantity(48T)</b>
Assay plate	96T	48T
Standard	5 x 1 mL	5 x 0.5 mL
HRP-conjugate	1 x 7 mL	1 x 3.5 mL
Antibody	1 x 7 mL	1 x 3.5 mL
TMB Substrate	1 x 12 mL	1 x 6 mL
Stop Solution	1 x 10 mL	1 x 5 mL
Sample Diluent	1 x 50 mL	1 x 25 mL
Wash Buffer(10x)	1 x 30 mL	1 x 15 mL
Adhesive Strip	4	4
Instruction Manual	1	1

## **STANDARD CONCENTRATION**

Standard	S1	S2	S3	S4	S5
Concentration ( $\mu\text{g}/\text{kg}$ )	0	0.15	0.45	1.35	4.05

## **STORAGE**

Unopened kit	Store at 2 - 8°C. Do not use the kit beyond the expiration date
Opened kit	May be stored for up to 1 month at 2 - 8° C.

**\*Provided this is within the expiration date of the kit.**

## **OTHER SUPPLIES REQUIRED**

- Microplate reader capable of measuring absorbance at 450 nm.
- An incubator which can provide stable incubation conditions up to 25°C
- Squirt bottle, manifold dispenser, or automated microplate washer.
- Centrifuge, Vortex mixer
- Analytical balance, 2 decimal place
- Single-channel micropipette(20  $\mu$ L-200  $\mu$ L, 100  $\mu$ L-1000  $\mu$ L)
- 300  $\mu$ L multichannel micropipette
- 50 mL and 500 mL graduated cylinders
- Deionized or distilled water
- Pipette and pipette tips
- Test tubes for dilution
- Ethyl alcohol absolute

## **PRECAUTIONS**

The Stop Solution provided with this kit is an acid solution. Wear eye, hand, face, and clothing protection when using this material.

### **Note:**

- Kindly use graduated containers to prepare the reagent.
- Bring all reagents to room temperature (20-25°C) before use for 30 min.
- Only the disposable tips can be used for the experiments and the tips must be changed when used for different reagents.
- Distilled water is recommended to be used to make the preparation for reagents. Contaminated water or container for reagent preparation will influence the detection result.

## **REAGENT PREPARATION**

- **40% Hydrous Ethanol:** Take 400 mL ethyl alcohol absolute into 600 mL deionized water, and shake well.
- **Wash Buffer (1x):** If crystals have formed in the concentrate, warm up to

room temperature and mix gently until the crystals have completely dissolved. Dilute 10 mL of **Wash Buffer (10x)** into 90 mL deionized or distilled water to prepare 100 mL of **Wash Buffer (1x)**. Keep it at 4°C for one month.

**Note:**

1. CUSABIO is only responsible for the kit itself, but not for the samples consumed during the assay. The user should calculate the possible amount of the samples used in the whole test. Please reserve sufficient samples in advance.
2. If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary.
3. Please predict the concentration before assaying. If values for these are not within the range of the standard curve, users must determine the optimal sample dilutions for their particular experiments.

## **SAMPLE COLLECTION AND STORAGE**

### **A. Feed samples**

1. Weigh  $5.00 \pm 0.05$ g smashed sample, put into 25mL **40% hydrous ethanol**, shake for 2 minutes.
2. Centrifuge at 4000 rpm/min for 5 min.
3. Take 500 $\mu$ L of supernatant into 500 $\mu$ L of **Sample Diluent** and shake well.
4. Take 50 $\mu$ L of sample for further analysis.  
Dilution factor of the samples: 10

### **STORAGE**

The prepared sample may be stored for up to one day at 2-8°C.

## ASSAY PROCEDURE

**Bring all reagents and samples to room temperature (20–25°C) before use. Centrifuge the sample again after thawing before the assay. It is recommended that all samples and standards be assayed in duplicate.**

1. Prepare all reagents and samples as directed in the previous sections.
2. Determine the number of wells to be used and put any remaining wells and the desiccant back into the pouch and seal the ziploc, store unused wells at 2–8°C.
3. Add 50  $\mu\text{L}$  of **Standard** or **Sample** per well. Then add 50  $\mu\text{L}$  of **HRP-conjugate** and 50  $\mu\text{L}$  of **Antibody** to each well. Cover the microtiter plate with a new adhesive strip and mix well, then incubate for 30 min at 25°C.
4. Aspirate each well and wash, repeating the process four times. Wash by filling each well with 250  $\mu\text{L}$  of **Wash Buffer (1x)** and let it stand for 30 seconds, complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining Wash Buffer by aspirating or decanting. Invert the plate and blot it against clean paper towels.
5. Add 100  $\mu\text{L}$  of **TMB Substrate** to each well, mix well. Incubate for 15 minutes at 25° C. Protect from light.
6. Add 50  $\mu\text{L}$  of **Stop Solution** to each well, gently tap the plate to ensure thorough mixing.
7. Determine the optical density of each well within 5 min, using a microplate reader set to 450nm (Recommend to read the OD value at the dual-wavelength: 450/630 nm within 5 min).

Note:

1. The final experimental results will be closely related to validity of the products, operation skills of the end users and the experimental environments.
2. Samples or reagents addition: Please carefully add samples to wells and mix gently to avoid foaming. Do not touch the well wall as possible. For each step in the procedure, total dispensing time for addition of reagents or samples to the assay plate should not exceed 10 min. This will ensure equal elapsed time for each pipetting step, without interruption. Duplication of all standards and specimens, although not required, is recommended. 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.
3. Incubation: To ensure accurate results, proper adhesion of plate sealers during incubation steps is necessary. Do not allow wells to sit uncovered for extended periods between incubation steps. Once reagents have been added to the well strips, DO NOT let the strips DRY at any time during the assay. Incubation time and temperature must be observed.
4. Washing: The wash procedure is critical. Complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining Wash Solution by aspirating or decanting and remove any drop of water and fingerprint on the bottom of the plate. Insufficient washing will result in poor precision and falsely elevated absorbance reading. 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.
5. Controlling of reaction time: Observe the change of color after adding Substrates (e.g. observation once every 10 min). Substrates should change from colorless or light blue to gradations of blue. If the color is too deep, add Stop Solution in advance to avoid excessively strong reaction which will result in inaccurate absorbance reading.
6. Substrates are easily contaminated. Substrates should remain colorless or light blue until added to the plate. Please protect it from light.
7. Stop Solution should be added to the plate in the same order as the Substrates. 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 Substrates.



## CALCULATION OF RESULTS

There are two methods to judge the results: the first one (A) is the rough judgment, while the second (B) is the quantitative determination. Note that the OD value of the sample has a negative correlation with Ochratoxin A in the sample.

### A:

Compare the sample average absorbance values with standards values, the Ochratoxin A concentration in the samples can be concluded. For example, the absorbance value of sample 1 is 0.799, the absorbance value of sample 2 is 1.329; absorbance values of standard are: 1.926, 1.579, 1.248, 0.711, 0.289, and the corresponding concentration are:  $0\mu\text{g}/\text{kg}$ ,  $0.15\mu\text{g}/\text{kg}$ ,  $0.45\mu\text{g}/\text{kg}$ ,  $1.35\mu\text{g}/\text{kg}$ ,  $4.05\mu\text{g}/\text{kg}$ ; then the Ochratoxin A in sample 1 and sample 2 are  $0.45\mu\text{g}/\text{kg}$  -  $1.35\mu\text{g}/\text{kg}$  and  $0.15\mu\text{g}/\text{kg}$  -  $0.45\mu\text{g}/\text{kg}$ ; Lastly the reader is multiplied by the corresponding Dilution factor of each sample and the actual concentration of sample is obtained.

### B:

The software offered together will facilitate the calculation process, it's suitable for accurate and fast analysis of large scale samples, please contact us.

### Note:

- Discard the substrate with any color that indicates the degeneration of this solution; when the absorbance value of S1 less than 0.5 indicates its degeneration.
- The optimum reaction temperature is  $25^{\circ}\text{C}$ , and too high or too low will result in the changes in the absorbance value and detecting sensitivity.