



User's Manual

THC ELISA Kit

REF DEIA-XYZ230

 96T



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This product is for research use only and is not intended for diagnostic use.

For illustrative purposes only. To perform the assay the instructions for use provided with the kit have to be used.

Creative Diagnostics

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PRODUCT INFORMATION

Intended Use

For the determination of trace quantities of THC metabolite and Parent THC in human urine, blood, or oral fluid.

General Description

Creative Diagnostics's THC ELISA Kit is a qualitative one-step kit designed for use as a screening device for the detection of drugs and/or their metabolites. The kit was designed for screening purposes and is intended for research use only. It is recommended that all suspect samples be confirmed by a quantitative method such as gas chromatography/mass spectrometry (GC/MS).

Principles of Testing

Creative Diagnostics's THC ELISA kit operates on the basis of competition between the drug or its metabolite in the sample and the drug-enzyme conjugate for a limited number of antibody binding sites. First, the sample or control is added to the microplate. Next, the drug-enzyme conjugate is added and the mixture is incubated at room temperature. During this incubation, the drug in the sample or the drug-enzyme conjugate binds to antibody immobilized in the microplate wells. After incubation, the plate is washed to remove any unbound sample or drug-enzyme conjugate. The presence of bound drug-enzyme conjugate is recognized by the addition of Substrate (TMB). After a 30 minute substrate incubation, the reaction is halted with the addition of an acid stop. The test can be read visually or with a microplate reader equipped with a 450 nm filter. The extent of color development is inversely proportional to the amount of drug in the sample or control. In other words, the absence of the drug in the sample will result in a dark yellow color, whereas the presence of the drug will result in light yellow to no color development.

Reagents And Materials Provided

1. **EIA Buffer:** 30 mL (ready-to-use). Phosphate buffered saline solution with bovine serum and a preservative. Provided for dilution of samples.
2. **Wash Buffer Concentrate (10x):** 20 mL. Phosphate buffered saline solution with a surfactant. Dilute 10 fold with deionized or ultrapure water before use. Diluted wash buffer is used to wash all unbound conjugate and samples from the plate after the conjugate incubation.
3. **Substrate:** 20 mL (ready-to-use). Stabilized 3, 3', 5, 5' Tetramethylbenzidine (TMB) plus Hydrogen Peroxide (H₂O₂) in a single bottle. It is used to develop the color in the wells after washing. Light sensitive.
4. **Drug-Enzyme Conjugate:** 14 mL (ready-to-use). Drug-horseradish peroxidase conjugate. Do not dilute.
5. **Antibody Coated Plate:** A 96 well Costar plate, in strips of 8 break-away wells, coated with anti-drug antiserum. The plate is ready for use as is. Do not wash.
6. **Acid Stop Solution:** 14 mL (ready-to-use). 1 N H₂SO₄ used to stop the enzyme reaction.
7. **Whole Blood Cut-off Calibrators: DO NOT DILUTE.**
 - a. Blood Negative Calibrator: 1 mL provided in a synthetic matrix (red cap).

- b. Blood Cut-off Calibrator-(THC-COOH): 1 mL provided in a synthetic matrix, 5 ng/mL (red cap).
8. **Urine Cut-off Calibrators: DO NOT DILUTE.**
- a. Urine Negative Calibrator: 1 mL provided in a synthetic matrix (yellow cap).
- b. Urine Cut-off Calibrator-(THC-COOH): 1 mL provided in a synthetic matrix, 50 ng/mL (yellow cap).

Materials Required But Not Supplied

1. Deionized water.
2. Precision pipettes that range from 10 µL - 1000 µL and disposable tips.
3. Graduated cylinder to dilute and mix wash buffer.
4. Plate cover or plastic film to cover plate during incubation.
5. Clean glassware (i.e. test tubes) to dilute samples.
6. Microplate reader with a 450 nm filter.

Storage

This kit can be used until the expiration date on the label when stored refrigerated at 2-8°C.

Specimen Collection And Preparation

Recommended minimum sample dilutions are listed below. These dilutions may change based on your laboratory's determination. All sample dilutions should be made in Creative Diagnostics's EIA Buffer.

- a. **Urine:** A dilution of 1:50 (i.e. 1 part sample to 49 parts provided EIA buffer) is required for optimal assay performance against the provided urine calibrators (yellow caps). Please contact Creative Diagnostics for assistance.
- b. **Whole blood:** A dilution of 1:5 (i.e. 1 part sample to 4 parts provided EIA Buffer) is required for optimal assay performance against the provided whole blood calibrators (red caps). Please contact Creative Diagnostics for assistance.
- c. **Oral Fluid:** No sample dilution is recommended for optimal assay performance. A centrifugation step may be used to spin down particulate matter. Please contact Creative Diagnostics for assistance.
- d. **Other sample types:** Please contact Creative Diagnostics for assistance.

Assay Procedure

The following test procedures can be run manually or on an automated instrument. Please contact Creative Diagnostics for assistance with protocols for automated instruments.

NOTE: Blood, Urine, and Oral Fluid Samples require different amounts of sample and conjugate be added. Follow the recommended sample amounts for each matrix listed below. For other sample types, please contact Creative Diagnostics.

1. Determine the number of wells to be used.
2. Gently mix the ready to use conjugate solution by inversion. Do not vortex. Store unused conjugate at 2-

8°C.

3. Choose the appropriate Creative Diagnostics calibrators to be used with your sample type. Do not dilute the calibrators provided by Creative Diagnostics.

4. **For Whole Blood Samples:**

Add 10 µL of sample, Creative Diagnostics calibrators or laboratory calibrators to the appropriate wells in duplicate. Add 100 µL of the ready to use drug-enzyme conjugate to each well. For manual runs use 8-channel pipette or 12-channel pipette for rapid addition.

For Urine Samples:

Add 10 µL of sample, Creative Diagnostics calibrators or laboratory calibrators to the appropriate wells in duplicate. Add 100 µL of the ready to use drug-enzyme conjugate to each well. For manual runs use 8-channel pipette or 12-channel pipette for rapid addition.

For Oral Fluid Samples:

Add 20 µL of sample or laboratory calibrators to the appropriate wells in duplicate. Add 100 µL of the ready to use drug-enzyme conjugate to each well. For manual runs use 8-channel pipette or 12-channel pipette for rapid addition.

5. For manual runs, mix by gently shaking plate. A microplate shaker may be used.
6. Cover plate with plastic film or plate cover and incubate at room temperature for 45 minutes.
7. During the conjugate incubation, dilute concentrated wash buffer 10 fold with deionized water (i.e. 20 mL of concentrated wash buffer plus 180 mL of deionized water). Mix thoroughly. Diluted wash buffer is stable for 5 days at room temperature or 7 days at 2-8°C.
8. Once the incubation is complete, dump or aspirate the liquid from the wells. Tap the plate on a clean lint-free towel to remove any remaining liquid in the wells.
9. Wash each well with 300 µL of diluted wash buffer. **Manual Wash:** For manual wash procedures repeat for a total of 3 washings, invert and tap dry the plate following each step. After completing the last wash step wipe the bottom of the wells with a lint-free towel to remove any liquid on the outside of the wells. **Automated Wash:** If an automated plate washer is used wash the plate for a total of 5 washings with 300 µL of diluted wash buffer. It is important for the automated washer to conduct a final aspirate cycle to eliminate residual amounts of wash buffer. Residual amounts of buffer in the wells will affect assay performance. Note: DI water should never be used for the plate wash.
10. Add 100 µL of the Substrate to each well. For manual runs, use a multi-channel pipetter for best results.
11. Incubate at room temperature for 30 minutes.
12. Add 100 µL of the Acid Stop (1N H₂SO₄) to each well to stop enzyme reaction. Mix gently before measuring absorbance. For automated systems a 10 second shake is sufficient. Measure the absorbance at a wavelength of 450 nm. Wells should be read within 2 hours of stopping the reaction.

NOTES

1. Desiccant bag must remain in foil pouch with unused strips. Keep zip lock pouch sealed when not in use to maintain a dry environment.
2. Use clean pipette tips for the buffer, drug-enzyme conjugate, controls and samples.
3. Before pipetting a reagent, rinse the pipette tip three times with that reagent.
4. When pipetting into the wells, **DO NOT** allow the pipette tip to touch the inside of the well or any of the reagent already inside the well. This may result in cross contamination.

- 5. Controls and samples should be assayed in duplicate.
- 6. Before substrate addition, wipe the outside bottom of the wells with a lint-free wiper to remove dust and fingerprints.
- 7. Gently mix specimens and reagents before use. Avoid vigorous agitation.

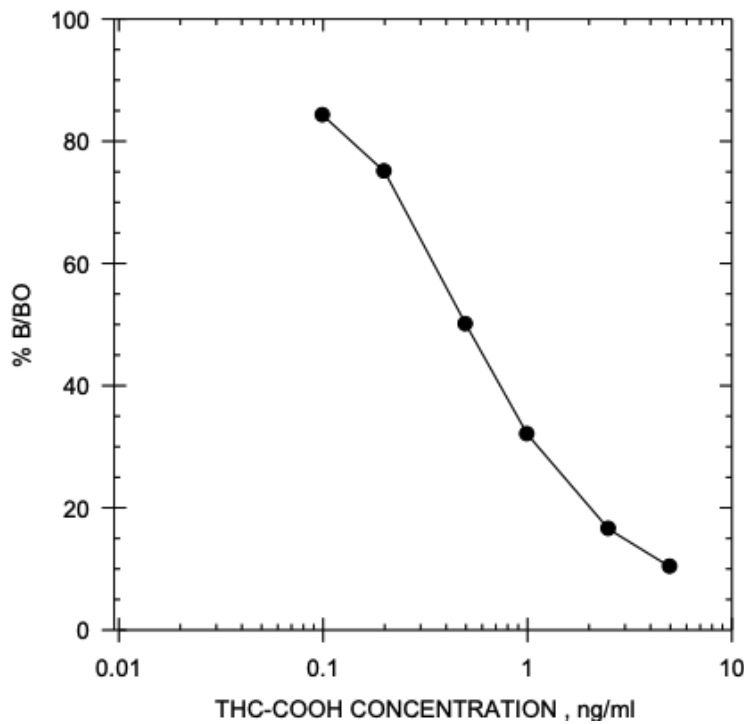
Interpretation Of Results

Positive Result: Samples with an absorbance less than or equal to the cutoff calibrator provided with the kit should be presumed positive. All positive samples should be confirmed by a quantitative method such as GC/MS.

Negative Result: Samples with an absorbance greater than the cutoff calibrator should be presumed negative.

Note: The kit was designed for screening purposes only. It is recommended that all suspect samples be confirmed by a quantitative method such as GC/MS or HPLC.

Typical Standard Curve



Sensitivity

Compound	I-50 in EIA Buffer
Δ^9 -THC-COOH	0.5 ng/mL

The term I-50 is used to define the sensitivity of the test. This number is derived from a standard curve generated with the drug. The drug concentration that shows 50% less color activity than the zero standard is considered to be the I-50.

Specificity

Compound	Compound Concentration (ng/mL)	THC-COOH Equivalents (ng/mL)	% Cross-Reactivity
Δ^9 -THC-COOH	0.5	0.5	100
Δ^8 -THC-COOH	0.6	0.5	88
11-OH- Δ^9 -THC	3.9	0.5	13
Δ^8 -THC	11	0.5	5
Δ^9 -THC	14	0.5	4
Cannabinol	24	0.5	2
Cannabidiol	4769	0.5	0.01

Note: THC-COOH equivalents represent 50% B/B₀ assay displacement in EIA Buffer.

The compounds having cross-reactivity below 0.01% did not show any significant reaction up to 10µg/mL

ALL THE FOLLOWING HAVE A CROSS-REACTIVITY <0.01%:

Acepromazine; Acetaminophen; Acetylsalicylic Acid; E-Amino-n-caproic Acid; Amitriptyline; Ascorbic Acid; Benzoic Acid; Caffeine; Chlordiazepoxide; Chlorpromazine; Clenbuterol; Codeine; Cotinine; Dexamethasone; Dextromethorphan; Diclofenac; Dimethyl Sulfoxide; Dipyrone; Doxepin; Ephedrine; Erythromycin; Ethyl p-aminobenzoate; Fenoprofen; Flunixin; Folic Acid; Folinic Acid; Furosemide; Gemfibrozil; Gentisic Acid; Glipizide; L-Glutamic Acid; Glutethimide; Glycopyrrolate; Heparin; Hippuric Acid; Hordenine; Hydrocortisone; Ibuprofen; Imipramine; Isoxsuprine; JWH-018; Lidocaine; Meperidine; Metaproterenol; Methadone; Methaqualone; Methocarbamol; Methylene Blue; 6- α Methylprednisolone; Nalorphine; Naproxen; Niacinamide; Nicotine; Nortriptyline; Olanzapine; Orphenidine; Oxyphenbutazone; Penicillin G-Potassium; Penicillin G-Procaine; Pentoxifylline; Phencyclidine; Phenothiazine; Phenylbutazone; Polyethylene Glycol; Prednisolone; Primidone; Procainamide; Procaine; Promazine; Pseudoephedrine; Pyrantel; Pyrilamine; Pyrimethamine; Quinidine; Quinine; Salbutamol; Salicylamide; Salvinorin A; Theophylline; Thiamine; Trimethoprim; Trimipramine; Uric Acid.

Precautions

- DO NOT** use kits or components beyond expiration date.
- DO NOT** mix conjugates and plates from different kit lots.
- DO NOT** pipette reagents by mouth.
- Pour Substrate out of the bottle into a clean reservoir. To prevent contamination of the substrate, **DO NOT** pipette out of the bottle.
- All specimens should be considered potentially infectious. Exercise proper handling precautions.
- Keep plate covered except when adding reagents, washing or reading.
- Kit components should be refrigerated at 2-8°C when not in use.
- Keep the cutoff calibrators stored at 2-8°C.
- Use aseptic technique when opening and removing reagents from vials and bottles.
- DO NOT** smoke, eat or drink in areas where specimens or reagents are being handled.
- Do not substitute DI water for the wash step of this protocol. Use only Creative Diagnostics's wash buffer.

12. Sodium Azide concentrations at 0.01% or less should not interfere with the assay provided that recommended dilutions are followed.
13. Do not reuse wells, they are for one use only.