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## BENZODIAZEPINE GROUP ULTRA

ELISA KIT INSTRUCTIONS PRODUCT #134319 & #134315 (RTU)  
FORENSIC USE ONLY

**INTENDED USE:** For the determination of trace quantities of Benzodiazepine and/or other metabolites in human urine, blood, and oral fluid.

### DESCRIPTION

Neogen Corporation's Benzodiazepine Group Ultra ELISA (Enzyme-Linked ImmunoSorbent Assay) test 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 forensic use only. It is recommended that all suspect samples be confirmed by a quantitative method such as gas chromatography/mass spectrometry (GC/MS).

### ASSAY PRINCIPLES

Neogen Corporation's test 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 3 times to remove any unbound sample or drug-enzyme conjugate. The presence of bound drug-enzyme conjugate is recognized by the addition of K-Blue® Substrate (TMB). After the 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.

### STORAGE AND STABILITY

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

### MATERIALS PROVIDED – SINGLE KIT (96 WELL)

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. **K-Blue Substrate:** 20 mL (ready-to-use). Stabilized 3,3',5,5' Tetramethylbenzidine (TMB) plus Hydrogen Peroxide (H<sub>2</sub>O<sub>2</sub>) 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 H<sub>2</sub>SO<sub>4</sub> 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- (Oxazepam): 1 mL provided in a synthetic matrix, 50 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- (Oxazepam): 1 mL provided in a synthetic matrix, 200 ng/mL (yellow cap).

**9. Oral Fluid Cut-off Calibrators: DO NOT DILUTE.**

- a. Oral Fluid Negative Calibrator: 1 mL provided in a synthetic matrix (clear cap).
- b. Oral Fluid Cut-off Calibrator- (Oxazepam): 1 mL provided in a synthetic matrix, 50 ng/mL (clear cap).

**MATERIALS PROVIDED – BULK KIT (480 WELL)**

1. **EIA Buffer:** 200 mL (ready-to-use). Phosphate buffered saline solution with bovine serum and a preservative. Provided for dilution of samples.
2. **Wash Buffer Concentrate (10X):** 100 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. **K-Blue Substrate:** 100 mL (ready-to-use). Stabilized 3, 3', 5, 5' Tetramethylbenzidine (TMB) plus Hydrogen Peroxide (H<sub>2</sub>O<sub>2</sub>) in a single bottle. It is used to develop the color in the wells after washing. Light sensitive.
4. **Drug-Enzyme Conjugate:** 5 x 14 mL (ready-to-use). Drug-horseradish peroxidase conjugate. Do not dilute.
5. **Antibody Coated Plate:** 5 x 96 well Costar plates, in strips of 8 break-away wells, coated with anti-drug antiserum. The plates are ready for use as is. Do not wash.
6. **Acid Stop Solution:** 90 mL (ready to use). 1 N H<sub>2</sub>SO<sub>4</sub> 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- (Oxazepam): 1 mL provided in a synthetic matrix, 50 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- (Oxazepam): 1 mL provided in a synthetic matrix, 200 ng/mL (yellow cap).
9. **Oral Fluid Cut-off Calibrators: DO NOT DILUTE.**
  - a. Oral Fluid Negative Calibrator: 1 mL provided in a synthetic matrix (clear cap).
  - b. Oral Fluid Cut-off Calibrator- (Oxazepam): 1 mL provided in a synthetic matrix, 50 ng/mL (clear cap).

**MATERIALS NEEDED BUT NOT PROVIDED**

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.

**OPTIONAL MATERIALS**

1. Microplate shaker.

**PRECAUTIONS AND NOTES**

1. **DO NOT** use kits or components beyond expiration date.
2. **DO NOT** mix conjugates and plates from different kit lots.
3. **DO NOT** pipette reagents by mouth.
4. Pour K-Blue Substrate out of the bottle into a clean reservoir. To prevent contamination of the substrate, **DO NOT** pipette out of the bottle.
5. All specimens should be considered potentially infectious. Exercise proper handling precautions.
6. Keep plate covered except when adding reagents, washing or reading.
7. Kit components should be refrigerated at all times when not in use.
8. Keep the cut-off calibrators stored at 2-8°C.
9. Use aseptic technique when opening and removing reagents from vials and bottles.
10. **DO NOT** smoke, eat or drink in areas where specimens or reagents are being handled.
11. Do not substitute DI water for the wash step of this protocol. Use only Neogen's wash buffer.
12. Sodium Azide concentrations at 0.01 % or less should not interfere with the assay.
13. Do not reuse wells, they are for one use only.

## PROCEDURAL NOTES

1. Desiccant bag must remain in zip lock bag 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, calibrators 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. Calibrators 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.

## SAMPLE TREATMENT

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

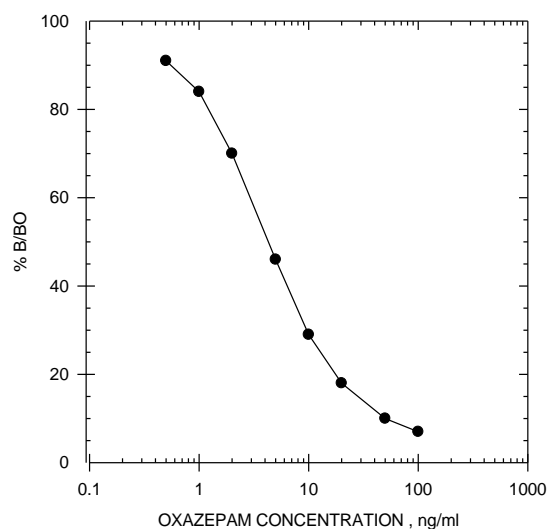
- a. **Urine:** A dilution of 1:10 (i.e. 1 part sample to 9 parts provided EIA buffer) is required for optimal assay performance against the provided urine calibrators (yellow caps). The assay has been optimized, however, it can be used for running sexual assault urine samples by modifying the sensitivity of the assay. Please contact your Neogen Representative 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 your Neogen Representative for assistance.
- c. **Oral Fluid:** 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 oral fluid calibrators (clear caps). A centrifugation step may be used to spin down the particulate matter prior to dilution. Please contact your Neogen Representative for assistance.
- d. **Other Forensic sample types:** Please contact your Neogen Representative for assistance.

## TEST PROCEDURES

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

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 Neogen calibrators to be used with your sample type. Do not dilute the calibrators provided by Neogen.
4. Add 10  $\mu\text{L}$  of sample, Neogen calibrators or laboratory calibrators to the appropriate wells in duplicate. Please contact your Neogen Representative with assistance concerning assay protocol and sample treatment changes.
5. Add 100  $\mu\text{L}$  of the ready to use drug-enzyme conjugate to each well. For manual runs use 8-channel pipetter or 12-channel pipetter for rapid addition.
6. For manual runs, mix by gently shaking plate. A microplate shaker may be used.
7. Cover plate with plastic film or plate cover and incubate at room temperature for 45 minutes.
8. 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.
9. 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.
10. Wash each well with 300  $\mu\text{L}$  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  $\mu\text{L}$  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.
11. Add 100  $\mu\text{L}$  of the K-Blue Substrate to each well. For manual runs, use a multi-channel pipetter for best results.
12. Incubate at room temperature for 30 minutes.
13. Add 100  $\mu\text{L}$  of the Acid Stop (1N  $\text{H}_2\text{SO}_4$ ) 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.

## STANDARD CURVE IN EIA BUFFER



## SENSITIVITY

Compound	I-50 in EIA Buffer
Oxazepam	6.30 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 in EIA Buffer. 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)	Oxazepam Equivalents (ng/mL)	% Cross-Reactivity
Oxazepam	6.3	6.30	100
Nordiazepam	3.5	6.30	179
Nimetazepam	4.0	6.30	159
Estazolam	4.1	6.30	154
Clobazam	4.3	6.30	146
Bromazepam	4.5	6.30	142
Diazepam	4.5	6.30	142
Etizolam	4.5	6.30	140
Pyrazolam	4.6	6.30	137
Clorazepate	4.8	6.30	132
Alphahydroxyalprazolam	4.8	6.30	132
Medazepam	4.9	6.30	129
Clonazolam	5.2	6.30	122
Flubromazepam	5.4	6.30	118
Nitrazepam	5.4	6.30	117
Triazolam	5.6	6.30	113
Alprazolam	5.7	6.30	112
Flualprazolam	5.7	6.30	112
Desalkylflurazepam	5.7	6.30	111
Lormetazepam	5.8	6.30	109
Flunitrazolam	5.9	6.30	107
Tetrazepam	6.1	6.30	104
Diclazepam	6.2	6.30	103
2-Hydroxyethylflurazepam	6.3	6.30	101

Flurazepam	6.3	6.30	99
Lorazepam	6.4	6.30	99
Alphahydroxytriazolam	6.5	6.30	97
Temazepam	6.6	6.30	96
N-Desmethylflunitrazepam	6.8	6.30	93
Clonazepam	6.8	6.30	93
Flubromazolam	6.9	6.30	92
Flunitrazepam	7.0	6.30	90
Delorazepam	7.1	6.30	90
Cloniprazepam	7.7	6.30	82
7-Aminonitrazepam	8.5	6.30	75
Prazepam	9.0	6.30	70
7-Aminoflunitrazepam	9.1	6.30	70
Halazepam	9.3	6.30	69
Chlordiazepoxide	9.3	6.30	68
Midazolam	10.3	6.30	62
7-Aminoclonazepam	10.6	6.30	60
Phenazepam	12.0	6.30	53
Meclonazepam	21.0	6.30	30
Nifoxipam	46.3	6.30	14
Ketazolam	97	6.30	7
Oxaprozin	298	6.30	2
Nefopam	4178	6.30	0.2

Note: Oxazepam equivalents represents 50% B/B<sub>0</sub> assay displacement in EIA Buffer.

*The compounds having cross-reactivity below 0.07% did not show any significant reaction up to 10µg/mL.*

**ALL THE FOLLOWING HAVE A CROSS-REACTIVITY <0.07%.**

Acetopromazine; Acetaminophen; Acetylsalicylic Acid; E-amino-n-caproic Acid; Amitriptyline; Ascorbic Acid; Benzoic Acid; Caffeine; Chlordiazepoxide; Chlorpromazine; Clenbuterol; Codeine; Cotinine; Dexamethasone; Dextromethorphan; Diclofenac; Dimethyl Sulfoxide; Doxepin; 1R, 1S(-) Ephedrine; 1S,2R(+) Ephedrine; Erythromycin; Ethyl p-amino benzoate; Fenoprofen; Flunixin; Folic Acid; Folinic Acid; Furosemide; Gemfibrozil; Gentic Acid; Glipizide; L-Glutamic Acid; Glutethimide; Glycopyrrolate; Heparin; Hippuric Acid; Hordenine; Hydrocortisone; Ibuprofen; Imipramine; Isoxsuprine; Lidocaine; Meperidine; Metaproterenol; Methadone; Methaqualone; Methocarbamol; Methylene Blue; Methylprednisolone; Mitragnine; Nalorphine; Naproxen; Niacinamide; Nicotine; Nortriptyline; Orphenadrine; Oxyphenbutazone; Penicillin G-Potassium; Penicillin G-Procaïne; Pentoxifylline; Phenothiazine; Phenylbutazone; Polyethylene Glycol; Prednisolone; Primidone; Procainamide; Procaine; Promazine; R,R(-) Pseudoephedrine; S, S(+) Pseudoephedrine; Pyrantel; Pyrimethamine; Pyrilamine; PCP; Quinidine; Quinine; Salbutamol; Salicylamide; Salicylic Acid; Salvinorin A; Sodium Azide; Theophylline; Thiamine; Trimethoprim; Trimipramine; Uric Acid; Zolpidem.

## RESULTS INTERPRETATION

**Positive Result:** Samples with an absorbance less than or equal to the cut-off calibrator 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 cut-off 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.

## TECHNICAL SUPPORT

For technical assistance, please contact our Technical Services Department at (859) 254-1221 or email at [techservice-toxicology@neogen.com](mailto:techservice-toxicology@neogen.com). Representatives are available Monday – Friday from 8:00 am – 6:00 pm EST.

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