Fluridone ELISA KIT

Cat. No.: DEIA6864
Pkg. Size: 100T

**Intended use**

The Fluridone ELISA KIT is suitable for the detection and quantitation of Fluridone in water (groundwater, surface water, well water).

**Principle Of The Test**

The Fluridone Kit applies the principles of enzyme linked immunosorbent assay (ELISA) to the determination of Fluridone. The sample to be tested is added, along with an enzyme conjugate, to a disposable test tube, followed by paramagnetic particles attached with antibodies specific to Fluridone. At this point a competitive reaction occurs between the Fluridone which may be in the sample and the enzyme labeled Fluridone analog for the antibody binding sites on the magnetic particles. The reaction is allowed to continue for twenty minutes. At the end of the incubation period, a magnetic field is applied to hold in the test tube the para-magnetic particles (with Fluridone and labeled Fluridone bound to the antibodies on the particles, in proportion to their original concentration), and allow the unbound reagents to be decanted. After decanting, the particles are washed with Washing Solution.

The presence of Fluridone is detected by adding the "Color Solution", which contains the enzyme substrate (hydrogen peroxide) and the chromogen (3,3',5,5'-tetramethylbenzidine). The enzyme-labeled Fluridone bound to the Fluridone antibody catalyzes the conversion of the substrate/chromogen mixture to a colored product. After an incubation period, the reaction is stopped and stabilized by the addition of a diluted acid (Stopping Solution). Since the labeled Fluridone (conjugate) was in competition with the unlabeled Fluridone (sample) for the antibody sites, the color developed is inversely proportional to the concentration of Fluridone in the sample.

**Reagents And Materials Provided**

1. Fluridone Antibody Coupled Paramagnetic Particles
   Fluridone antibody (rabbit anti-Fluridone) covalently bound to paramagnetic particles suspended in a buffered solution with preservative and stabilizers.
   100 test kit: one 65 mL vial
2. Fluridone Enzyme Conjugate
   Horseradish peroxidase (HRP) labeled Fluridone analog diluted in a buffered solution with additive, preservative and stabilizers.
   100 test kit: one 35 mL vial
3. Fluridone Standards
   Four concentrations (0.5, 2.0, 7.5, 15.0 ppb) of Fluridone standards in distilled water with preservative and stabilizers. Each vial contains 2.0 mL.
4. Control
   A concentration (approximately 6.0 ppb) of Fluridone in distilled water with preservative and stabilizers. A 2.0 mL volume is supplied in one vial.
5. Diluent/Zero Standard
   Distilled water with preservative and stabilizers without any detectable Fluridone.
   100 test kit: one 35 mL vial
6. Color Solution
A solution of hydrogen peroxide and 3,3',5,5'-tetramethylbenzidine in an organic base.
100 test kit: one 60 mL vial

7. Stopping Solution
A solution of diluted acid.
100 test kit: one 60 mL vial

8. Washing Solution T
Preserved deionized water with preservative and detergent.
100 test kit: one 250 mL vial

9. Test Tubes
Glass tubes are packaged in a box.
100 test kit: three 36 tube boxes

Materials Required But Not Supplied

1. Pipets* Precision pipets capable of delivering 50, 250 and 500 µL and a 1.0 mL repeating pipet.
2. Vortex Mixer* Thermolyne Maxi Mix, Scientific Industries Vortex Genie, or equivalent
3. Magnetic Separation System*
4. Photometer* capable of readings at 450 nm
* Please contact us for supplier information

Storage
Store all reagents at 2-8°C. Do not freeze. Reagents may be used until the expiration date on the box. The test tubes and Washing Solution require no special storage condition and may be stored separately from the reagents to conserve refrigerator space.
Consult state, local, and federal regulations for proper disposal of all reagents.

Specimen Collection And Handling

This procedure is recommended for use with water samples. Other samples may require modifications to the procedure and should be thoroughly validated.
Samples containing gross particulate matter should be filtered (e.g. Uniprep 0.45 µm, Whatman, Inc.) to remove particles.
Samples which have been preserved with monochloroacetic acid or other acids, should be neutralized with strong base e.g. 6N NaOH, prior to assay.
If the Fluridone concentration of a sample exceeds 15.0 ppb, the sample is subject to repeat testing using a diluted sample. A ten-fold or greater dilution of the sample is recommended with an appropriate amount of Diluent/Zero Standard or Sample Diluent. For example, in a separate test tube make a ten-fold dilution by adding 100 uL of the sample to 900 uL of Diluent/Zero Standard. Mix thoroughly before assaying. Perform the assay according to the Assay Procedure and obtain final results by multiplying the value obtained by the dilution factor (e.g. 10).

Reagent Preparation
All reagents must be allowed to come to room temperature. The antibody coupled paramagnetic particles should be mixed thoroughly before use.

Assay Steps

1. Label test tubes for standards, control, and samples.
Table 1.

<table>
<thead>
<tr>
<th>Tube Number</th>
<th>Contents of Tube</th>
</tr>
</thead>
<tbody>
<tr>
<td>1,2</td>
<td>Diluent/Zero Standard, 0 ppb</td>
</tr>
<tr>
<td>3,4</td>
<td>Standard 1, 0.5 ppb</td>
</tr>
<tr>
<td>5,6</td>
<td>Standard 2, 2.0 ppb</td>
</tr>
<tr>
<td>7,8</td>
<td>Standard 3, 7.5 ppb</td>
</tr>
<tr>
<td>9,10</td>
<td>Standard 4, 15.0 ppb</td>
</tr>
<tr>
<td>11,12</td>
<td>Control</td>
</tr>
<tr>
<td>13, 14</td>
<td>Sample 1</td>
</tr>
<tr>
<td>15,16</td>
<td>Sample 2</td>
</tr>
<tr>
<td>17, 18</td>
<td>Sample 3</td>
</tr>
</tbody>
</table>

2. Add 150 µL of the appropriate standard, control, or sample.
3. Add 250 µL of Fluridone Enzyme Conjugate to each tube.
4. Mix the Fluridone Antibody Coupled Paramagnetic Particles thoroughly and add 500 µL to each tube.
5. Vortex for 1 to 2 seconds minimizing foaming.
6. Incubate for 20 minutes at room temperature.
7. Separate in the Magnetic Separation System for two minutes.
8. Decant and gently blot all tubes briefly in a consistent manner.
9. Add 1 mL of Washing Solution T to each tube and allow them to remain in the magnetic separation unit for two minutes.
10. Decant and gently blot all tubes briefly in a consistent manner.
11. Repeat Steps 9 and 10 two additional times, for a total of 3 washes.
12. Remove the rack from the separator and add 500 µL of Color Solution to each tube.
13. Vortex for 1 to 2 seconds minimizing foaming.
14. Incubate for 20 minutes at room temperature.
15. Add 500 µL of Stopping Solution to each tube.
16. Add 1 mL Washing Solution to a clean test tube. Use as blank in Step 17.
17. Read results at 450 nm within 15 minutes after adding the Stopping Solution.

**Quality Control**

A control solution at approximately 6.0 ppb of Fluridone is provided with the Fluridone Assay kit. It is recommended that it be included in every run and treated in the same manner as unknown samples. Acceptable limits should be established by each laboratory.
**Calculation**

**Manual Calculations**
1. Calculate the mean absorbance value for each of the standards.
2. Calculate the %B/Bo for each standard by dividing the mean absorbance value for the standard by the mean absorbance value for the Diluent/Zero Standard.
3. Construct a standard curve by plotting the %B/Bo for each standard on vertical logit (Y) axis versus the corresponding Fluridone concentration on horizontal logarithmic (X) axis on the graph paper provided.
4. %B/Bo for controls and samples will then yield levels in ppb of Fluridone by interpolation using the standard curve.

**Photometric Analyzer**
Some instrument manufacturers make available photometers allowing for calibration curves to be automatically calculated and stored. Refer to instrument operating manual for detailed instructions. To obtain results for the Fluridone Assay on instruments allowing data tranformation the following parameter settings are recommended:
- Data Reduct: Lin. Regression
- Xformation: Ln/Logit
- Read Mode: Absorbance
- Wavelength: 450 nm
- Units: PPB
- # Rgt Blk: 0
- Calibrators:
  - # of Cals: 5
  - # of Reps: 2
- Concentrations:
  - #1: 0.00 PPB
  - #2: 0.5 PPB
  - #3: 2.0 PPB
  - #4: 7.5 PPB
  - #5: 15.0 PPB
- Range: 0.50 - 15.0
- Correlation: 0.990
- Rep. %CV: 10%

**Reference Values**
In a study with water samples from locations across the U.S., the Fluridone Assay was shown to correlate well with HPLC ($r^2 = 0.985$).

**Sensitivity**
The Fluridone Assay has an estimated minimum detectable concentration, based on 4 SD from zero = 0.24 ppb.

**Specificity**
The cross-reactivity of the Fluridone Assay for various analogues can be expressed as the least detectable dose (LDD) which is estimated at 90% B/Bo, or as the dose required for 50% absorbance inhibition (50% B/Bo).

Table 2.
The following compounds demonstrated no reactivity in the Fluridone Assay at concentrations up to 1000 ppb: aldicarb, aldicarb sulfoxide, aldicarb sulfone, atrazine, ametryn, benomyl, butylate, captan, carbaryl, carbendazim, carbofuran, cyanazine, 2,4-D, 1,3-dichloropropene, dinoseb, MCPA, metribuzin, pentachlorophenol, picloram, propazine, simazine, terbufos, thiabendazole, thiophanate-methyl, triclopyr, and trifularin.

**Recovery**

Four groundwater samples were spiked with various levels of Fluridone and then assayed using the Fluridone Assay. The following results were obtained:

Table 3:
Reproducibility

<table>
<thead>
<tr>
<th>Amount of Fluridone Added (ppb)</th>
<th>Recovery</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean (ppb)</td>
</tr>
<tr>
<td>1.0</td>
<td>1.09</td>
</tr>
<tr>
<td>2.0</td>
<td>2.27</td>
</tr>
<tr>
<td>4.0</td>
<td>4.51</td>
</tr>
<tr>
<td>8.0</td>
<td>8.68</td>
</tr>
<tr>
<td>12.0</td>
<td>12.0</td>
</tr>
<tr>
<td>Average</td>
<td></td>
</tr>
</tbody>
</table>

Precautions

1. As with all immunoassays, a consistent technique is the key to optimal performance. To obtain the greatest precision, be sure to treat each tube in an identical manner.
2. Add reagents directly to the bottom of the tube while avoiding contact between the reagents and the pipet tip. This will help assure consistent quantities of reagent in the test mixture.
3. Avoid cross-contaminations and carryover of reagents by using clean pipets for each sample addition and by avoiding contact between reagent droplets on the tubes and pipet tips.
4. Avoid excessive foam formation during vortexing.
5. The magnetic separation system consists of two parts: an upper rack which will securely hold the test tubes and a lower separator which contains the magnets used to attract the antibody coupled paramagnetic particles. During incubations the upper rack is removed from the lower separator so that the paramagnetic particles remain suspended during the incubation. For separation steps, the rack and the separator are combined to pull the paramagnetic particles to the sides of the tubes.
6. To obtain optimum assay precision, it is important to perform the separation steps carefully and consistently. Decant the rack by slowly inverting away from the operator using a smooth and continuous turning action so the liquid flows consistently along only one side of the test tube. While still inverted, place the rack on an absorbent pad and allow to drain. Lifting the rack and replacing gently onto the pad several times will ensure complete removal of the liquid from the rim of the tube. Do not bang the rack.
7. Mix the antibody coupled paramagnetic particles just prior to pipetting.
8. Do not use any reagents beyond their stated shelf life.
9. Avoid contact of Stopping Solution (diluted sulfuric acid) with skin and mucous membranes. If this reagent comes in contact with skin, wash with water.
Limitations

The Fluridone Assay will detect Fluridone and related pyridazinone compounds to different degrees. Refer to specificity table for data on several of the compounds. The Fluridone Assay kit provides screening results. As with any analytical technique (GC, HPLC, etc...) positive results requiring some action should be confirmed by an alternative method.

The total time required for pipetting the magnetic particles should be kept to two minutes or less, therefore the total number of tubes that can be assayed in a run should be adjusted accordingly.