

Metolachlor

• Intended Use

For the detection and quantitation of Metolachlor and related acetanilides in water (groundwater, surface water, well water). For soil, crop, and food use contact the company for application bulletins and/or specific matrix validation guidelines.

• Principle

The Abraxis Metolachlor Kit applies the principles of enzyme linked immunosorbent assay (ELISA) to the determination of Metolachlor and related acetanilides. 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 metolachlor. At this point a competitive reaction occurs between the Metolachlor or other acetanilides which may be in the sample and the enzyme labeled Metolachlor analog for the antibody binding sites on the magnetic particles. The reaction is allowed to continue for thirty (30) minutes. At the end of the incubation period, a magnetic field is applied to hold in the test tube the paramagnetic particles (with Metolachlor and labeled Metolachlor 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 Metolachlor 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 Metolachlor bound to the Metolachlor 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 Metolachlor (conjugate) was in competition with the unlabeled Metolachlor (sample) for the antibody sites, the color developed is inversely proportional to the concentration of Metolachlor in the sample.

• Reagents

The Abraxis Metolachlor Kit contains the following items:

1. Metolachlor Antibody Coupled Paramagnetic Particles

Metolachlor antibody (rabbit anti-Metolachlor) covalently bound to paramagnetic particles suspended in a buffered solution with preservative and stabilizers.

100 test kit: one 65 mL vial

2. Metolachlor Enzyme Conjugate

Horse radish peroxidase (HRP) labeled Metolachlor analog diluted in a buffered solution with preservative and stabilizers.

100 test kit: one 35 mL vial

3. Metolachlor Standards

Three concentrations (0.1, 1.0, 5.0 ppb) of Metolachlor standards in distilled water with preservative and stabilizers. Each vial contains 2.0 mL.

4. Control

A concentration (approximately 2 ppb) of Metolachlor 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 Metolachlor.

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 65 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 detergent.

100 test kit: one 250 mL vial

9. Test Tubes

Polystyrene tubes (22) are packaged in a box.

100 test kit: three 36 tube boxes

• Reagent Storage and Stability

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.

• Materials Required but Not Provided

In addition to the reagents provided, the following items are essential for the performance of the test:

Pipets* Precision pipets capable of delivering 200, and 500 μ L and a 1.0 mL repeating pipet.

Vortex Mixer* Thermolyne Maxi Mix, Scientific Industries Vortex Genie, or equivalent

Magnetic Separation System*

Photometer* capable of readings at 450 nm

* Please contact Abraxis for supplier information.

• Sample Information

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. 0.2 μ m Anotop™ 25 Plus, 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 Metolachlor concentration of a sample exceeds 5 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 μ L of the sample to 900 μ L 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.

• Procedural Notes and Precautions

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.

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.

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.

Avoid foam formation during vortexing.

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.

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 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.

Mix the antibody coupled paramagnetic particles just prior to pipetting.

Do not use any reagents beyond their stated shelf life.

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 Abraxis Metolachlor Assay will detect Metolachlor and related acetanilides to different degrees. Refer to specificity table for data on several of the acetanilides. The Abraxis Metolachlor 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 (2) minutes or less, therefore the total number of tubes that can be assayed in a run should be adjusted accordingly.

• Quality Control

A control solution at approximately 2 ppb of Metolachlor is provided with the Abraxis Metolachlor 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.

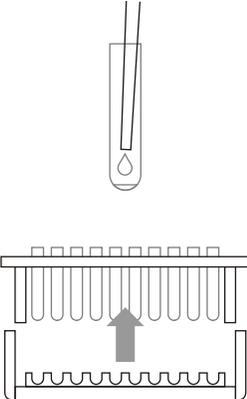
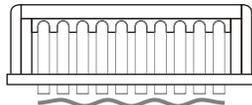
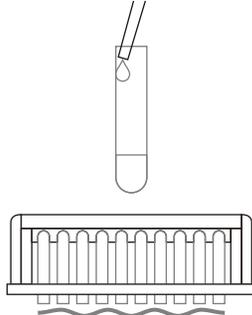
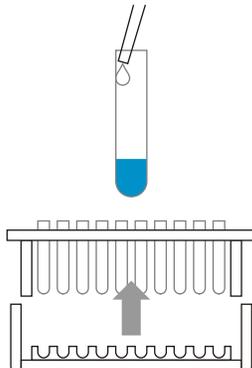
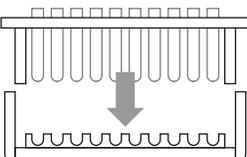
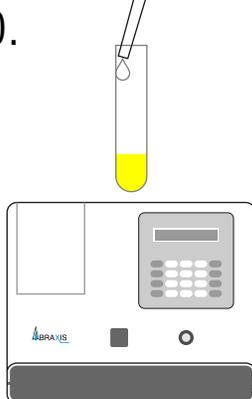
• Assay Procedure

Read Reagent Preparation, Procedural Notes and Precautions before proceeding.

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

Tube Number	Contents of Tube
1,2	Diluent/Zero Standard, 0 ppb
3,4	Standard 1, 0.1 ppb
5,6	Standard 2, 1.0 ppb
7,8	Standard 3, 5.0 ppb
9	Control
10	Sample 1
11	Sample 2
12	Sample 3

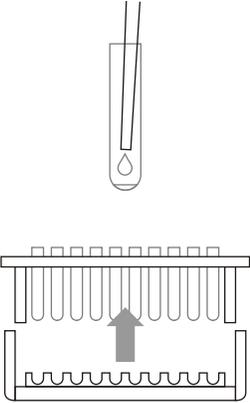
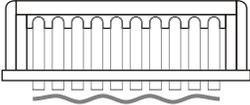
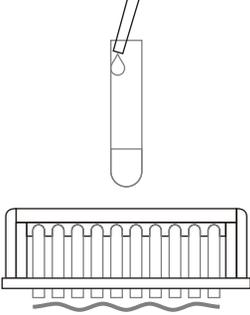
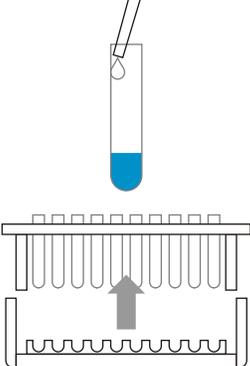
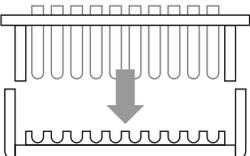
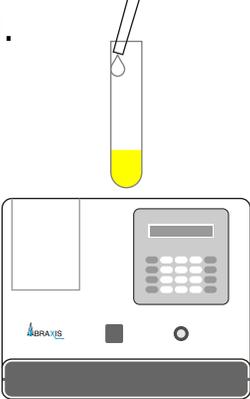
METOLACHLOR DETAILED FLOWCHART

<p>1.</p>  <p><i>Remove</i> upper rack from magnetic base. Label test tubes for Standards, Control, and Samples.</p> <table border="1"> <thead> <tr> <th>Tube #</th> <th>Content</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, 0.1 ppb</td> </tr> <tr> <td>5, 6</td> <td>Standard 2, 1.0 ppb</td> </tr> <tr> <td>7, 8</td> <td>Standard 3, 5.0 ppb</td> </tr> <tr> <td>9</td> <td>Control</td> </tr> <tr> <td>10</td> <td>Sample 1</td> </tr> <tr> <td>11</td> <td>Sample 2</td> </tr> <tr> <td>12</td> <td>Sample 3</td> </tr> </tbody> </table> <p>Add 200 μL without of either Standards, Control or Samples to the bottom of each test tube by inserting the pipette tip all the way into the bottom of the tube without touching the sides of the tube.</p>	Tube #	Content	1, 2	Diluent/Zero Standard 0 ppb	3, 4	Standard 1, 0.1 ppb	5, 6	Standard 2, 1.0 ppb	7, 8	Standard 3, 5.0 ppb	9	Control	10	Sample 1	11	Sample 2	12	Sample 3	<p>6.</p>  <p>Do not separate upper rack from lower base. Using a smooth motion, <i>invert</i> the combined rack assembly over a sink and pour out the tube contents; keep inverted and gently blot the test tube rims on several layers of paper toweling.</p>
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1, 2	Diluent/Zero Standard 0 ppb																		
3, 4	Standard 1, 0.1 ppb																		
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11	Sample 2																		
12	Sample 3																		
<p>2.</p>  <p>Add 250 μL of Metolachlor Enzyme Conjugate down the inside wall of each tube by aiming the pipet tip 1/4" to 1/2" below the tube rim without touching the rim or tube wall with the pipet tip; deliver liquid gently.</p>	<p>7.</p>  <p>Add 1 mL of Washing Solution down the inside wall of each tube by using the technique described in Box 2. <i>Wait 2 minutes</i>. Using a smooth motion, invert the combined rack assembly over a sink and pour out the tube contents: keep inverted and gently blot the test tube rims on several layers of paper toweling. Repeat this step.</p>																		
<p>3.</p>  <p>Add 500 μL of thoroughly mixed Metolachlor Antibody Coupled Magnetic Particles down the inside wall of each tube by using the technique described in Box 2. <i>Vortex</i> for 1 to 2 seconds (at low speed to minimize foaming).</p>	<p>8.</p>  <p><i>Lift</i> the upper rack (with its tubes) off the magnetic base; add 500 μL of Color Reagent down the inside wall of each tube by using the technique described in Box 2. <i>Vortex</i> for 1 to 2 seconds (at low Speed to minimize foaming).</p>																		
<p>4.</p>  <p><i>React</i> 30 minutes at room temperature (15° - 30°C).</p>	<p>9.</p>  <p><i>React</i> for 20 minutes at room temperature (15° - 30° C). During this period, add 1 mL of Washing Solution into a clean tube for use as an instrument blank in Step 10.</p>																		
<p>5.</p>  <p><i>Combine</i> the upper rack with the magnetic base; press all tubes into base; allow 2 minutes for the particles to separate.</p>	<p>10.</p>  <p>Add 500 μL of Stopping Solution down the inside wall of each tube by using the technique previously described. <i>Read</i> results at 450 nm within 15 minutes after adding the Stopping Solution. <i>Multiply</i> results of samples by the appropriate dilution factor (if any).</p> <p>[Safety Caution: Stopping Solution contains diluted sulfuric acid.]</p>																		

For Ordering or Technical Assistance Contact:
 ABRAXIS, LLC 54 Steamwhistle Drive, Warminster, PA 18974
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Metolachlor Magnetic Particle Kit Part # 500061, 100 Test

METOLACHLOR CONCISE FLOWCHART

<p>1.</p>  <p>Separate the rack.</p> <p>Add 200 μL of either Standards, Control or Samples to the bottom of each test tube.</p>	<p>6.</p>  <p>Invert the combined rack.</p> <p>Blot gently.</p>
<p>2.</p>  <p>Add 250 μL of Metolachlor Enzyme Conjugate to each test tube.</p>	<p>7.</p>  <p>Add 1 mL of Washing Solution.</p> <p>Wait 2 minutes.</p> <p>Invert the combined rack.</p> <p>Blot gently.</p> <p>Repeat this step.</p>
<p>3.</p>  <p>Add 500 μL of mixed Magnetic Particles to each test tube.</p> <p>Vortex.</p>	<p>8.</p>  <p>Separate the rack.</p> <p>Add 500 μL of Color Reagent to each test tube.</p> <p>Vortex.</p>
<p>4.</p>  <p>Incubate for 30 minutes.</p>	<p>9.</p>  <p>Incubate for 20 minutes.</p> <p>Prepare blank.</p>
<p>5.</p>  <p>Combine the rack and magnetic base.</p> <p>Seat all tubes.</p> <p>Wait 2 minutes.</p>	<p>10.</p>  <p>Add 500 μL of Stopping Solution to each test tube.</p> <p>Read OD 450</p>

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