

METOLACHLOR in Water

Immunoassay Method*

Scope and Application: For water

This method analyzes for Metolachlor in water. Sample calibrators and reagents are added to cuvettes coated with Metolachlor-specific antibodies. The color that develops is then measured and compared with the color measurements of the calibrators. The test requires about 30 minutes for complete analysis. As many as 20 cuvettes (18 samples and 2 calibrators) can be run simultaneously.

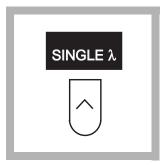
Tips and Techniques

- Read the entire procedure before starting. Identify and have ready all the necessary reagents, cuvettes, and other apparatus before beginning the analysis.
- Timing is critical; follow instructions carefully.
- A consistent technique when mixing the cuvettes is critical to this test. The best results come from using the cuvette rack and mixing as described in *Using the 1-cm MicroCuvette Rack*. Cuvettes can be mixed individually, but test results may not be as consistent.
- Handle the cuvettes carefully. Scratches on the inside or outside may cause erroneous results. Carefully clean the outside
 of the cuvettes with a clean absorbent cloth or tissue before placing them into the instrument. (Kimwipe[®] tissues are
 provided with the kit.)
- Antibody cuvettes and enzyme conjugate are made in matched lots. Do not mix reagent lots.
- To avoid damaging the Color Developing Solution, do not expose it to direct sunlight.
- The cuvette rack is designed to be inverted with the cuvettes in place. This is especially helpful when running many samples at once; the cuvettes can remain in the rack and be processed together until they are read in the Immunoassay Pocket Colorimeter.
- Twenty Antibody Cuvettes are provided with each reagent set. One Antibody Cuvette will be used for each calibrator and each sample. Cuvettes are not reusable.
- Ensure the 1cm MicroCell adapter is installed in the DR/4000.

Note: Hach Company recommends wearing protective nitrile gloves for this procedure.

^{*} This test is semi-quantitative. Results are expressed as greater or less than the threshold value used.

Immunoassay



1. Press the soft key under **SINGLE** λ .

Press the soft key under **GO TO** λ .

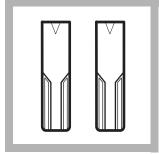
Select 450 nm by pressing the numeric keys **450**.

Press: **ENTER**

Note: The Flow Cell and Sipper Modules cannot be used with this procedure.

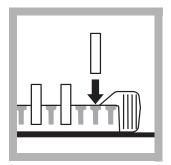


2. The display will show: **ZERO REQUIRED**

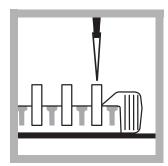


3. Label an Antibody Cuvette for each calibrator and each sample to be tested.

Note: As many as 20 cuvettes may be tested at one time and may comprise any combination of samples and calibrators.

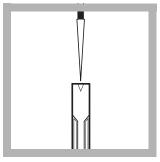


4. Place the cuvettes into the rack snugly.



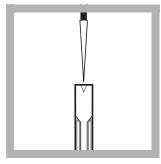
5. Pipet 0.5 mL of each calibrator into the appropriately labeled cuvette.

Note: Use a new pipette tip for each sample.



6. Pipet 0.5 mL of each sample to be tested into the appropriately labeled cuvette.

Note: Use a new pipette tip for each sample.



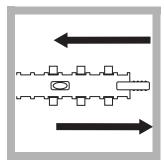
7. Immediately pipet 0.5 mL of Metolachlor Enzyme Conjugate into each cuvette.



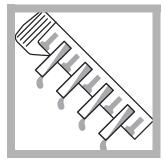
8. Key **2000**. Press the soft key under **START TIMER**.

A 20-minute reaction time will begin.

Immediately mix the contents of the cuvettes for 30 seconds using the technique described in *Using the 1-cm MicroCuvette Rack*.



9. After 10 minutes mix the contents of the rack for 30 seconds using the technique described in "Using the 1-cm MicroCuvette Rack" on page 5.



10. At the end of the 20-minute period, discard the contents of all the cuvettes into an appropriate waste container.

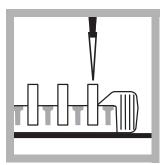


11. Wash each cuvette forcefully and thoroughly four times with deionized water. Empty the rinse water into the waste container.

Note: Ensure most of the water is drained from the cuvettes by turning the cuvettes upside down and tapping them lightly on a paper towel

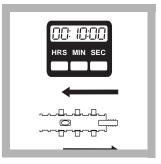
Color Development

Note: Timing is critical; follow instructions carefully.



12. With the cuvettes still held snugly in the rack, pipet 0.5 mL of Color Developing Solution into each Antibody Cuvette.

Note: Use a new pipette tip for each cuvette.



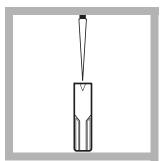
13. Key **1000**. Press the soft key under **START TIMER**.

A reaction period will begin. Mix following the instructions in *Using the 1-cm MicroCuvette Rack*.



14. After 5 minutes, mix the contents of the rack a second time for a period of 30 seconds using the same technique.

Note: Solutions will turn blue in some or all of the cuvettes.



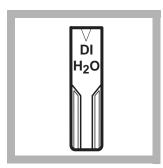
15. At the end of the 10-minute reaction period, pipette 0.5 mL of Stop Solution into each cuvette in the same order as the Color Developing Solution was added in *step 12*.

Slide the rack for 20 seconds using the technique described in *Using the 1-cm MicroCuvette Rack*.

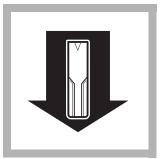
Note: Blue solutions will turn yellow with the addition of the Stop Solution.

Note: The same pipette tip can be used repeatedly for this step.

Measuring the Color



16. Label and fill a Zeroing Cuvette with deionized water. Wipe the outside of all the cuvettes with a tissue to remove water, smudges, and fingerprints.



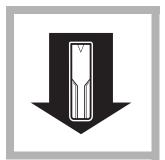
17. Place the filled zeroing cuvette into the cell holder with the arrow pointing left.

Orient the arrow in the same direction for all cuvettes.



18. Press the soft key under **ZERO**.

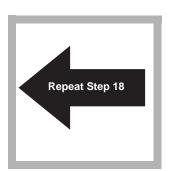
The display will show: **0.000 Abs**



19. Place the first calibrator into the cell holder. Read the results.

The display will give an absorbance reading. Record the results for each calibrator and sample.

Note: See the Instrument Manual for more information on taking a reading.



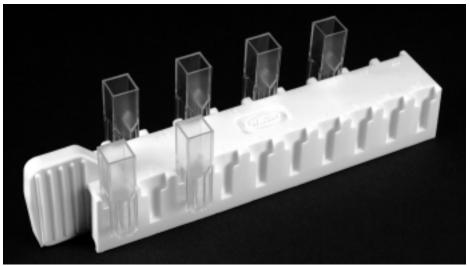
20. Repeat *step 19* for all remaining calibrators and samples.

See *Interpreting and Reporting Results* for help with interpretation of results.

Using the 1-cm MicroCuvette Rack

This rack (see *Figure 1*) has been designed specifically to aid in achieving precise and accurate results when using the immunoassay technique to analyze several samples at the same time.

Figure 1 The 1-cm MicroCuvette Rack



Loading the Rack — The cuvette rack is designed so that it may be inverted with the cuvettes in place. Identify each cuvette with a sample or calibrator number and place all the cuvettes in the rack before beginning the procedure. Fit the cuvettes snugly into the rack, but do not force them or they may be difficult to remove and their contents may spill. The cuvettes should remain in place when the rack is inverted and tapped lightly.

Mixing — Set the rack on a hard, flat surface that is at least twice the length of the rack. Hold the rack by one end and vigorously slide it back and forth along its long axis for 30 seconds. The rack should move through a distance equal to its own length in each direction.

Interpreting and Reporting Results

There is an inverse relationship between the concentration of Metolachlor and the reading. In other words, the higher the reading, the lower the concentration of Metolachlor.

If the sample reading is	the sample Metolachlor Concentration is	
less than calibrator reading	greater than the calibrator concentration	
greater than calibrator reading	less than the calibrator concentration	

Example

Readings:

0.5 ppm Metolachlor Calibrator: **0.475 ABS**

2.0 ppm Metolachlor Calibrator: **0.245 ABS**

Sample #1: 0.140 ABS

Sample #2: 0.300 ABS

Sample #3: **0.550 ABS**

Interpretation

Sample #1 — Sample reading is less than the readings for both calibrators. Therefore the sample concentration of Metolachlor is greater than both 0.5 ppb and 2.0 ppb Metolachlor.

Sample #2 — Sample reading is between the readings for the 0.5 ppb and 2.0 ppb Metolachlor calibrators. Therefore the sample concentration of Metolachlor is between 0.5 ppb and 2.0 ppb.

Sample #3 — Sample reading is greater than the readings for both calibrators. Therefore the sample concentration of Metolachlor is less than both 2.0 ppb and 0.5 ppb.

Storing and Handling Reagents

- Wear protective gloves and eyewear.
- When storing reagent sets for extended periods of time, keep them out of direct sunlight. Store reagents at a temperature of 4 °C when not in use.
- Keep the foil pouch containing the Metolachlor Antibody Cuvettes sealed when not in use.
- If Stop Solution comes in contact with eyes, wash thoroughly for 15 minutes with cold water and seek immediate medical help.

Sensitivity

The Metolachlor immunoassay cannot differentiate between the various triazines and metabolites, but it detects their presence in differing degrees.

Table 1 Required Concentrations for Selected Chemicals

Compound	Concentration to give a positive result at 0.5 ppb Metolachlor	Concentration to give a positive result at 2.0 ppb Metolachlor	
Acetochlor	74 ppm	398 ppm	
Butachlor	84 ppb	550 ppb	
2 Chloro-2'6'- Diethylacetaniline	8 ppm	60 ppm	
2,6-Diethylaniline	61 ppm	313 ppm	
Propachlor	60 ppb	295 ppb	

The following compounds are not detectable at 10,000 ppb.

Atrazine Carbofuran Carbendazim

Aldicarb 2,4-D

Diazoton Chlorpyirfos

Sample Collection and Storage

Collect samples in a clean glass bottle. Do not pre-rinse the bottle with the sample. If the sample cannot be analyzed immediately, store the sample at 4 °C. Samples may be kept for as long as 14 days. Warm the samples to room temperature before analysis.

Summary of Method

Hach immunoassay tests use antigen/antibody reactions to test for specific organic compounds in water and soil. Metolachlor-specific antibodies, attached to the walls of plastic cuvettes, selectively bind and remove Metolachlor from complex sample matrices. A prepared sample and a reagent containing enzyme-conjugate molecules (analyte molecules attached to molecules of an enzyme) are added to the Antibody Cuvettes. During incubation, enzyme-conjugate molecules and Metolachlor compete for binding sites on the antibodies. Samples with higher levels of analyte will have more antibody sites occupied by Metolachlor and fewer antibody sites occupied by the enzyme-conjugate molecules.

After incubation, the sample and unbound enzyme conjugate are washed from the cuvette and a color-development reagent is added. The enzyme in the conjugate catalyzes the development of color. Therefore, there is an inverse relationship between color intensity and the amount of Metolachlor in the sample. The resulting color is then compared with a calibrator to determine whether the Metolachlor concentration in the sample is greater or less than the threshold levels.

Required Reagents Description Reagent Set, Metolachlor*	U nit 20 cuvettes	Cat. No28135-00
Required Apparatus		
Caps, flip spout	2/pkg	25818-02
Cell Adapter, 1-cm MicroCell	each	48588-00
Marker, laboratory	each	20920-00
Rack, for 1-cm Micro Cuvettes	each	48799-00
Wipes, disposable	box	20970-00
TenSette®, Pipet, 0.1–1.0 mL	each	19700-01
Tips, for pipettor 19700-01	1000/pkg	21856-28

METOLACHLOR in Water

^{*} Immunoassay components are manufactured for Hach Company by Beacon Analytical Systems, Inc.

