

Atrazine

• Intended Use

For the detection and quantitation of atrazine and related triazines 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 Atrazine Kit applies the principles of enzyme linked immunosorbent assay (ELISA) to the determination of atrazine and related triazines. 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 triazines. At this point a competitive reaction occurs between the triazine which may be in the sample and the enzyme labeled atrazine for the antibody binding sites on the magnetic particles. The reaction is allowed to continue for fifteen (15) minutes. At the end of the incubation period, a magnetic field is applied to hold in the test tube the para-magnetic particles (with atrazine and labeled atrazine 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 atrazine 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 atrazine bound to the atrazine 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 atrazine (conjugate) was in competition with the unlabeled atrazine (sample) for the antibody sites, **the color developed is inversely proportional to the concentration of atrazine in the sample.**

• Reagents

The Abraxis Atrazine Kit contains the following items:

1. Atrazine Antibody Coupled Paramagnetic Particles

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

100 test kit: one 65 mL vial

2. Atrazine Enzyme Conjugate

Horseradish peroxidase (HRP) labeled atrazine analog diluted in a buffered solution with preservative and stabilizers.

100 test kit: one 35 mL vial

3. Atrazine Standards

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

4. Control

A concentration (approximately 3 ppb) of atrazine 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 atrazine.

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

Preserved deionized water.

100 test kit: one 250 mL vial

9. Test Tubes

Polystyrene tubes (36) 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 250, 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 atrazine 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 Atrazine Assay will detect atrazine and related triazines to different degrees. Refer to specificity table for data on several of the triazines. The Abraxis Atrazine 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 3 ppb of atrazine is provided with the Abraxis Atrazine 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

2. Add 200 or 250 uL of the appropriate standard, control, or sample.
3. Add 250 uL of Atrazine Enzyme Conjugate to each tube.
4. Mix the Atrazine Antibody Coupled Paramagnetic Particles thoroughly and add 500 uL to each tube.
5. Vortex for 1 to 2 seconds minimizing foaming.
6. Incubate for 15 minutes at room temperature.
7. Separate in the Magnetic Separation System for **two (2) minutes**.
8. Decant and **gently** blot all tubes briefly in a consistent manner.
9. Add 1 mL of Washing Solution to each tube and allow them to remain in the magnetic separation unit for **two (2) minutes**.
10. Decant and **gently** blot all tubes briefly in a consistent manner.
11. Repeat Steps 9 and 10 an additional time.
12. Remove the rack from the separator and add 500 uL 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 uL 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.

• Results

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 atrazine 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 atrazine by interpolation using the standard curve.
(Contact Abraxis for detailed application information on specific photometers.)

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 Abraxis Atrazine Assay on instruments allowing data transformation the following parameter settings are recommended:

Data Reduct : Lin. Regression
 Xformation : Ln/LogitB
 Read Mode : Absorbance
 Wavelength : 450 nm
 Units : PPB
 # Rgt Blk : 0

Calibrators:
 # of Cals : 4
 # of Reps : 2

Concentrations:
 #1: 0.00 PPB
 #2: 0.10 PPB
 #3: 1.00 PPB
 #4: 5.00 PPB

Range : 0.05 - 5.00
 Correlation : 0.990
 Rep. %CV : 10%

• Expected Results

In a study with water samples from locations across the U.S., the Abraxis Atrazine Assay was shown to correlate well with another commercial Atrazine immunoassay (r = 0.971).

• Performance Data

Precision

The following results were obtained:

Control	1	2	3
Replicates	5	5	5
Days	5	5	5
n	25	25	25
Mean (ppb)	1.34	2.65	3.99
% CV (within assay)	6.6	7.0	7.9
% CV (between assay)	2.9	3.0	5.1

Sensitivity

The Abraxis Atrazine Assay has an estimated minimum detectable concentration, based on a 90% B/Bo of 50 ppt.

Recovery

Five (5) groundwater samples, were spiked with various levels of atrazine and then assayed using the Abraxis Atrazine Assay. The following results were obtained:

Amount of Atrazine Added (ppb)	Recovery -----		
	Mean (ppb)	S.D. (ppb)	%
0.5	0.55	0.09	110
1.0	1.09	0.15	109
2.0	2.16	0.14	108
4.0	3.92	0.27	98
Average			106

Specificity

The cross-reactivity of the Abraxis Atrazine Assay for various triazine 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).

Compound	LDD (ppb)	50% B/Bo (ppb)
Atrazine	0.050	0.70
Propazine	0.084	1.18
Ametryn	0.022	0.44
Prometryn	0.052	0.80
Prometon	0.140	2.20
Desethyl Atrazine	0.250	4.75
Terbutryn	0.340	210
Simazine	0.760	3.40
Desisopropyl Atrazine	29	970
Cyanazine	0.800	47
2-Hydroxy Atrazine	0.960	20

The following compounds demonstrated no reactivity in the Abraxis Atrazine Assay at concentrations up to 1000 ppb: aldicarb, aldicarb sulfoxide, aldicarb sulfone, alachlor, benomyl, butachlor, butylate, captan, carbaryl, carbendazim, carbofuran, 2,4-D, 1,3-dichloropropene, dinoseb, MCPA, metolachlor, metribuzin, pentachlorophenol, picloram, propachlor, terbufos, thiabendazole, and thiophanate-methyl.

• Ordering information

Abraxis Atrazine Assay Kit 100T PN 500001
 Sample Diluent PN 500002
 Standard Set PN 500003

• Assistance

For ordering or technical assistance contact:

Abraxis LLC
 Sales Department
 54 Steamwhistle Drive
 Warminster, Pennsylvania, 18974

Phone: (215) 357-3911 * Fax: (215) 357-5232

Email: info@abraxiskits.com

WEB: www.abraxiskits.com

• General Limited Warranty

Abraxis LLC warrants the products manufactured by the Company, against defects and workmanship when used in accordance with the applicable instructions for a period not to extend beyond the product's printed expiration date. **Abraxis makes no other warranty, expressed or implied. There is no warranty of merchantability or fitness for a particular purpose.**

300001

R122605

NOTE: This ELISA kit has been verified under the EPA Environmental Testing Verification Program (March, 2004 report).



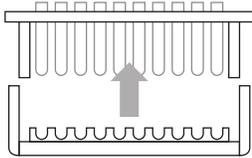
ATRAZINE DETAILED FLOWCHART

1.



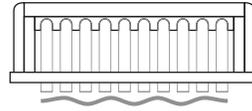
Remove upper rack from magnetic base. Label test tubes for Standards, Control, and Samples.

Tube #	Content
1,2	Diluent/Zero, 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



Add 200 or 250 μ L 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.

6.

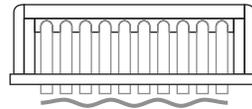


Do not separate upper rack from lower base. Using a smooth motion, *invert* the combined rack assembly over a sink and pour out the tube contents; keep inverted and **gently blot** the test tube rims on several layers of paper toweling.

7.



Add 1 mL of Washing Solution down the inside wall of each tube by using the technique described in Box 2. Wait 2 minutes. 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.



2.

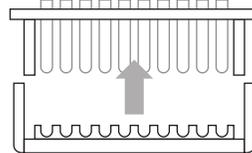


Add 250 μ L of Atrazine Enzyme Conjugate down the inside wall of each tube by using the technique described in Box 2. Vortex for 1 to 2 seconds (at low speed to minimize foaming).

8.



Lift the upper rack (with its tubes) off the magnetic base; add 500 μ L of Color Reagent down the inside wall of each tube by using the technique described in Box 2. Vortex for 1 to 2 seconds (at low speed to minimize foaming).



3.



Add 500 μ L of thoroughly mixed Atrazine Antibody Coupled Magnetic Particles down the inside wall of each tube by using the technique described in Box 2. Vortex for 1 to 2 seconds (at low speed to minimize foaming).

9.



Incubate 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.

4.

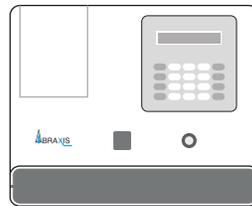


Incubate 15 minutes at room temperature (15°- 30°C).

10.



Add 500 μ L of Stopping Solution down the inside wall of each tube by using the technique previously described. Read results at 450 nm within 15 minutes after adding the Stopping Solution. Multiply results of samples by the appropriate dilution factor (if any).



[Safety Caution: Stopping Solution contains diluted sulfuric acid.]

For Ordering or Technical Assistance Contact:

ABRAXIS, LLC 54 Steamwhistle Drive, Warminster, PA 18974
215-357-3911 Fax: 215-357-5232

Atrazine Magnetic Particle Kit Part # 500001, 100 Test

ATRAZINE CONCISE FLOWCHART

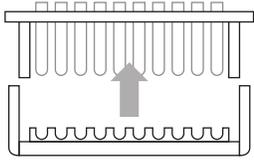
1.



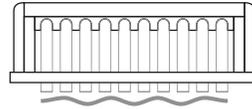
Separate the rack.

Add 200 or 250 μL of either Standards, Control or Samples to the bottom of each test tube.

NOTE: Same chosen volume should be added for standards and samples.



6.



Invert the combined rack.

Blot **gently**.

7.



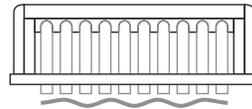
Add 1 mL of Washing Solution.

Wait 2 minutes.

Invert the combined rack.

Blot **gently**.

Repeat this step.



2.



Add 250 μL of mixed Atrazine Enzyme Conjugate to each test tube.

Vortex.

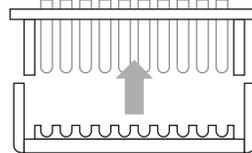
8.



Separate the rack.

Add 500 μL of Color Reagent to each test tube.

Vortex.



3.



Add 500 μL of mixed Magnetic Particles to each test tube.

Vortex.

9.



Incubate for 20 minutes.

Prepare blank.

4.



Incubate for 15 minutes.

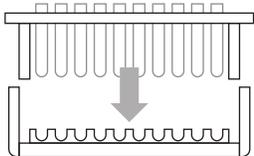
10.



Add 500 μL of Stopping Solution to each test tube.

Read OD 450

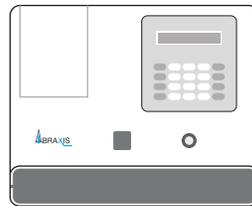
5.



Combine the rack and magnetic base.

Seat all tubes.

Wait 2 minutes.



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Phone: 215-357-3911 Fax: 215-357-5232

Atrazine Magnetic Particle Kit Part # 500001, 100 Test