Method number: PV2006

Target Concentration: 0.2 mg/m³ (ACGIH TLV-TWA)

Procedure: Samples are collected by drawing known volumes of air through impingers containing water. Samples are analyzed by high performance liquid chromatography (HPLC) using a UV detector.

Recommended air volume and sampling rate: 60 minutes at 1 Lpm (60 L)

Detection limit of the overall procedure: 0.004 mg/m³ (Based on the recommended air volume.)

Status of method: Partially Validated method. This method has been only partially evaluated and is presented for information and trial use.

December 1987 (final) Yihlin Chan

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1 General discussion

1.1 Background

The OSHA Analytical Laboratory received a set of samples requesting analysis of amitrole. The samples had been collected in impingers containing water. This report describes the preliminary validation of the sampling method and the analytical method developed for amitrole.

1.2 Toxic effects (This section is for information only and should not be taken as the basis for OSHA policy.)

"The oral LD$_{50}$ of amitrole in mice was 11,000 or 14,700 and was 25,000 mg/kg in rats. The intravenous LD$_{50}$ was 5000 mg/kg in mice. The dermal application of 10,000 mg/kg was tolerated by rats and rabbits without any systemic effects. The exposure of rats for four hours to an aerosol of 439 mg/m$^3$ did not produce symptoms of poisoning and did not irritate the eyes and the upper respiratory tract. It has a very slight irritating effect on the skin and eyes of rabbits." (Ref. 5.5)

The ACGIH TLV Committee recommended the removal of the A2 designation as a suspected human carcinogen from the listing for amitrole in 1984. (Ref. 5.5)

1.3 Potential workplace exposure (Ref. 5.4)

Amitrole is used as an herbicide and a plant growth regulator. No estimate on the extent of workplace exposure could be found.

1.4 Physical properties (Ref. 5.1 and 5.2)

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chemical name</td>
<td>3-Amino-1,2,4-triazole</td>
</tr>
<tr>
<td>Synonyms</td>
<td>3-Amino-1H-1,2,4-triazole; aminotriazole; ATA; ENT 25445; Amizol; Cytrol; Weedazol</td>
</tr>
<tr>
<td>CAS no.</td>
<td>61-82-5</td>
</tr>
<tr>
<td>Molecular formula</td>
<td>C$_2$H$_4$N$_4$</td>
</tr>
<tr>
<td>Molecular weight</td>
<td>84.1</td>
</tr>
<tr>
<td>Appearance</td>
<td>Colorless crystals; bitter taste</td>
</tr>
<tr>
<td>Melting point</td>
<td>159 °C</td>
</tr>
<tr>
<td>Solubility</td>
<td>Soluble in water (28% at 25 °C) and ethanol (26% at 75 °C); slightly soluble in chloroform, methylene chloride, acetonitrile, and ethyl acetate; insoluble in ether, acetone, and hydrocarbons</td>
</tr>
<tr>
<td>UV scan</td>
<td>See Figure 3.</td>
</tr>
</tbody>
</table>

Structure:

1.5 Detection limit of the analytical procedure

The detection limit of the analytical procedure is 0.51 ng per injection. This is the amount of analyte which will give a peak whose height is approximately 5 times the baseline noise.
2 Sampling procedure

2.1 Apparatus and reagents

2.1.1 A personal sampling pump that can be calibrated to within ±5% of the recommended flow rate when attached to the impinger.

2.1.2 Impinger

2.1.3 Distilled water

2.2 Sampling procedure (Ref. 5.3)

2.2.1 Calibrate pump.

2.2.2 Attach the collection device to the shirt collar or within the breathing zone. Position the tubing so as not to interfere with the work of the employee.

2.2.3 Turn on pump and record the starting time.

2.2.4 Check the pump flow periodically.

2.2.5 Prepare a blank.

2.2.6 At the end of the sampling period, turn off the pump and record the ending time.

2.2.7 Transfer the impinger solution to a vial. Rinse the impinger with a small amount of water and pour the rinse solution into the vial. Wrap the vial end-to-end with a Form OSHA-21 seal.

2.3 Recommended air volume and sampling rate

2.3.1 The recommended air volume is 60 L.

2.3.2 The recommended sampling rate is 1 Lpm.

2.4 Extraction efficiency

Not performed. Samples are analyzed directly.

2.5 Retention efficiency

Three impingers containing 15 mL of water were each spiked with 12.71 µg of amitrole. Humid air (80% RH, 60 L @ 1 Lpm) was drawn through the solutions. The solutions were each made to 15.0 mL with water and analyzed. The average recovery was 87.5%.

<table>
<thead>
<tr>
<th>table 2.5</th>
<th>amitrole retention efficiency</th>
</tr>
</thead>
<tbody>
<tr>
<td>sample</td>
<td>µg spiked</td>
</tr>
<tr>
<td>#</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>12.71</td>
</tr>
<tr>
<td>2</td>
<td>12.71</td>
</tr>
<tr>
<td>3</td>
<td>12.71</td>
</tr>
<tr>
<td>average</td>
<td></td>
</tr>
</tbody>
</table>
2.6 Storage

Three impingers containing 15 mL of water were each spiked with 12.71 µg of amitrole. Humid air (80% RH, 60 L @ 1 Lpm) was drawn through the solutions. The solutions were transferred to scintillation vials and stored at room temperature in the dark for 33 days. The average recovery was 79.9%.

<table>
<thead>
<tr>
<th>sample #</th>
<th>µg spiked</th>
<th>µg recovered</th>
<th>% recovered</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>12.71</td>
<td>9.34</td>
<td>73.5</td>
</tr>
<tr>
<td>2</td>
<td>12.71</td>
<td>10.48</td>
<td>82.4</td>
</tr>
<tr>
<td>3</td>
<td>12.71</td>
<td>10.62</td>
<td>83.6</td>
</tr>
</tbody>
</table>

average = 79.8%

2.7 Interferences

There are no known interferences to the sampling procedure.

3 Analytical method

3.1 Apparatus

3.1.1 High performance liquid chromatograph

3.1.2 Chromasil C18 column or equivalent

3.1.3 UV detector

3.1.4 Strip chart recorder

3.2 Reagents

3.2.1 Water, HPLC grade

3.2.2 Amitrole, EPA #0200

3.3 Standard preparation

Weigh 5 to 10 mg of amitrole in a 10-mL volumetric flask. Add water to the mark. Make a series of standards by diluting bulk with water to a suitable working range.

3.4 Sample preparation

The volumes of the samples were made to 15.0 mL with distilled water.

3.5 Analysis

3.5.1 Instrument conditions

<table>
<thead>
<tr>
<th>Column:</th>
<th>Chromasil C18</th>
</tr>
</thead>
<tbody>
<tr>
<td>Eluent:</td>
<td>100% water</td>
</tr>
<tr>
<td>Flow rate:</td>
<td>1.0 mL/min</td>
</tr>
</tbody>
</table>
Detector: 205 nm
Injection size: 30 uL
Retention time: 6.0 min

3.5.2 Chromatogram:

![Chromatogram of Amitrole](image)

Figure 1. Chromatogram of Amitrole

3.6 Interferences

3.6.1 Any collected compound that has the same retention time as amitrole and absorbs at 205 nm is a potential interference. Generally, chromatographic conditions can be altered to separate an interference from the analyte.

3.6.2 Retention time alone on a single column is not proof of a chemical identity. Confirmation by other means should be sought whenever possible.

3.7 Calculations

3.7.1 A calibration curve is constructed by plotting standard concentrations versus detector response (see Figure 2).

3.7.2 The concentration of a sample is determined from the calibration curve.

3.7.3 The air concentration is determined by the formula:

\[
\text{mg} / \text{m}^3 = \left( \frac{\mu g}{mL} \right) \left( \frac{15 mL}{\text{air volume, L}} \right)
\]

4 Recommendations for further study

4.1 Investigate a more convenient sampling device, such as a sorbent tube or a filter.

4.2 Develop a fully validated method.
5 References


