## TIER I HUMAN HEALTH NONCANCER CRITERIA

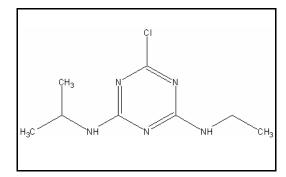
## **ATRAZINE**

CAS RN: 77-47-4 Water Solubility: 3.4 mg/L  $Log K_{ow}$ : 4.645

Reference Dose: 0.007 mg/kg/day

Carcinogenicity Weight-of-

Evidence Classification: Class D; Not Classifiable



#### Standard

The human health noncancer atrazine criterion for drinking water sources is 920  $\mu$ g/L. The human health noncancer criterion for nondrinking water sources is 15,000  $\mu$ g/L.

#### **Calculations**

BAF predicted based on Log  $K_{ow}$ Log  $K_{ow} = 2.396$  (CLOGP program),  $K_{ow} = 248.9$ Trophic level 3 FCM = 1.010; trophic level 4 FCM = 1.002

$$f_{\rm fd} = 1/(1 + (0.00000024~kg/L)(K_{ow})) = 1.0$$

Baseline BAF<sub>T3</sub> = 
$$(FCM)(K_{ow}) = (248.9)(1.010) = 253.9$$

Baseline BAF<sub>T4</sub> = 
$$(248.9)(1.002) = 249.4$$

Human health BAF<sub>T3</sub> = [(253.9)(0.0182)+1](1.0) = 5.621

Human health BAF<sub>T4</sub> = 
$$[(249.4)(0.0310)+1](1.0) = 8.731$$

## Acceptable Daily Exposure:

From the IRIS database:

Critical Effect: Decreased body weight gain

$$ADE = \frac{NOAEL}{UF} = \frac{3.5 \text{ mg/kg-day}}{100} = 0.035 \text{ mg/kg/d}$$

#### Calculation of Criteria:

Non Drinking Water HNV = 
$$[(0.035)(70)(0.8)]/0.01+[(0.0036)(5.621)+(0.0114)(8.731)]$$
  
= 15,000 µg/L

Drinking Water HNV = 
$$[(0.035)(70)(0.8)]/2+[(0.0036)(5.621)+(0.0114)(8.731)]$$
  
= 920 µg/L

#### References

- 1. USEPA 1993. Integrated Risk Information System (IRIS database) chemical file for atrazine (CAS # 1912-24-9).
- 2. Leo,A. and D.Weininger 1997. Daylight Software CLogP Version 3.15+ for Unix Pomona Medical Chemistry Project, Pomona College, Claremont, CA. Distributed by Daylight Chemical Information Systems, Inc., 3952 Claremont St., Irving, CA 92714 (Reference for the Log K<sub>ow</sub>)

## Acronyms

ADE	Acceptable Daily Exposure
BAF	Bioaccumulation Factor
CAS RN	Chemical Abstract Service Registry Number
FCM	Food Chain Multiplier
IRIS	Integrated Risk Information System
K <sub>ow</sub>	Octanol-Water Partition Coefficient
LOAEL	Lowest observed adverse effect level
NOAEL	No observed adverse effect level
P (superscript)	Predicted value
RPLC	Reverse-phase Liquid Chromatography
UF	Uncertainty factor

# **Revision History**

August 13, 1997 - Criteria first developed March 24, 2000 - Criteria rechecked (no modifications). Fact sheet updated.

# **Contact Information**

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