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Data Evaluation Report on the Acute Toxicity of Thidiazuron Technical to Freshwater Invertebrates - Daphnia

PMRA Submission Number {.......}

EPA MRID Number 46203503

Data Requirement:

PMRA DATA CODE

EPA DP Barcode

OECD Data Point EPA MRID

D294536

46203503

EPA Guideline

§72-2

Test material:

Thidiazuron Technical

Common name:

Thidiazuron

Chemical name:

IUPAC: Not reported

CAS name: Not reported CAS No.: 51707-55-2 Synonyms: AE B049537 Purity: 99.5% (w:w)

Primary Reviewer: Greg Hess

Staff Scientist, Dynamac Corporation

Signature:

Date: 4/1/04

QC Reviewer: Christie E. Padova Staff Scientist, Dynamac Corporation

Signature: Date: 4/22/04

Primary Reviewer: Bill Evans Biologist

OPP/EFED/ERB - I

Date: 5/26/04

Secondary Reviewer(s):

{EPA/OECD/PMRA}

Date:

Reference/Submission No.:

Company Code: **Active Code:** 

EPA PC Code: 120301

**Date Evaluation Completed:** 

CITATION: Blankinship, A.S., et al. 2003. Thidiazuron: A 48-Hour Static Acute Toxicity Test with the Cladoceran (Daphnia magna). Unpublished study performed by Wildlife International, Ltd., Easton, MD. Laboratory Project No. 149A-151. Study sponsored by Bayer CropScience, Frankfurt am Main, Germany. Study initiated February 18, 2003 and completed May 7, 2003.



#### **EXECUTIVE SUMMARY:**

The 48-hour acute toxicity of Thidiazuron Technical to the Cladoceran, *Daphnia magna*, was studied under static conditions. Neonate (<24-hour old) daphnids were exposed to the test material at nominal concentrations of 0 (negative control), 0.31, 0.63, 1.3, 2.5, 5.0, or 10 ppm. Mean-measured concentrations were <0.201 (<LOQ, control), 0.30, 0.60, 1.2, 2.3, 4.7, and 9.5 ppm a.i.

By 48 hours, mortality/immobility was 5% in the control group, 0% in the 0.30 through 2.3 ppm a.i. groups, 25% in the 4.7 ppm a.i. group, and 100% in the 9.5 ppm a.i. group. The 48-hour LC<sub>50</sub>/EC<sub>50</sub> (with 95% C.I.) was 5.7 (4.7-9.5) ppm a.i., which categorizes Thidiazuron Technical as moderately toxic to the water flea (*Daphnia magna*) on an acute toxicity basis. Lethargy was observed in 100% of surviving daphnids from the 9.5 ppm a.i. group after 24 hours, and in 5 and 40% of surviving daphnids from the 2.3 and 4.7 ppm a.i. groups, respectively, after 48 hours (100% mortality was observed at 9.5 ppm a.i. by 48 hours). The 48-hour NOEC and LOEC values were 1.2 and 2.3 ppm a.i., based on sub-lethal effects data (most sensitive endpoint).

This study is scientifically sound and satisfies the guideline requirements for an acute toxicity study with freshwater invertebrates (§72-2). This study is classified as CORE.

# Results Synopsis

Test Organism Age (eg. 1<sup>st</sup> instar): Neonates, <24 hours old Test Type (Flow-through, Static, Static Renewal): Static

### 48-Hour

LC<sub>50</sub>/EC<sub>50</sub>: 5.7 ppm a.i.

95% C.I.: 4.7-9.5 ppm a.i.

NOEC: 1.2 ppm a.i. (based on sub-lethal effects)

LOEC: 2.3 ppm a.i.

Endpoints affected: Mortality/immobility and sub-lethal effects

Most sensitive endpoint: Sub-lethal effects

### I. MATERIALS AND METHODS

**GUIDELINE FOLLOWED:** 

The study protocol was based on procedures outlined in the OECD Guideline No. 202 (1984); the U.S. EPA OPPTS No. 850.1010 (Draft, 1996); and ASTM Standard E729-88a (1994). Deviations from U.S. EPA §72-2 included:

- 1. Pre-test mortality of the laboratory culture and/or brood was not described.
- 2. The biomass loading rate was not specified.
- 3. The water hardness (136 mg/L as CaCO<sub>3</sub>) was nearly three times higher than recommended (40-48 mg/L as CaCO<sub>3</sub>).
- 4. The pH range (8.4-8.7) was greater than recommended (7.2-7.6).
- 5. Aeration of the test vessels was not addressed.

These deviations did not affect the acceptability or validity of the study.

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**COMPLIANCE:** 

Signed and dated GLP, Quality Assurance, and Data Confidentiality statements were provided. This study was conducted in accordance with GLP standards of the U.S. EPA (40 CFR Part 160 and 192), OECD (ENV/MC/CHEM (98)17), and Japan MAFF (11 NohSan, Notification No. 6283, Agricultural Production Bureau, 1 October 1999; p. 3).

### A. MATERIALS:

1. Test Material

Thidiazuron Technical

Description:

Powder

Lot No./Batch No.:

107623-03 (Product code: AE B049537 00 1D99 0003)

Purity:

99.5% (w:w) a.i.

Stability of Compound Under Test Conditions:

The stability of the test substance in the dilution water during the course of the study was demonstrated by analytical determination at 0 and 48 hours. Recoveries were 91.9-96.8% of nominal concentrations at 0 hours and 90.2-98.6% of nominal at 48 hours (Table 1, p. 17).

Storage conditions of test chemicals:

Stored at room temperature.

OECD requires water solubility, stability in water and light,  $pK_{a}$ ,  $P_{ow}$  and vapor pressure of the test compound. OECD requirements were not reported.

### 2. Test organism:

Species:

Daphnia magna

Age at test initiation:

Neonates, <24 hours old

Source:

In-house laboratory cultures; neonates were obtained from five individual adult daphnids.

# **B. STUDY DESIGN:**

# 1. Experimental Conditions

- a. Range-finding Study: The definitive nominal test concentrations were selected in consultation with the sponsor, and were based upon the results of an exploratory range-finding toxicity test. The results of the range-finding study were not reported (p. 9).
- b. Definitive Study

Table 1: Experimental Parameters

		Remarks Criteria		
Parameter	Details			
Acclimation period:	Continuous laboratory cultures were maintained (at least 13 days).			
Conditions: (same as test or not)	Same as test			
Feeding:	Daphnia cultures were fed a mixture of yeast, Cerophyll and trout chow with a suspension of the freshwater green alga, Selenastrum capricornutum.	EPA requires 7 day minimum acclimation period.		
Health: (any mortality observed)	No signs of disease or stress.	.2. 8		
Duration of the test	48 hours	EPA requires 48 hours		
Test condition - static/flow through	Static			
Type of dilution system (for flow through method)  Renewal rate (for static renewal)	N/A N/A	EPA requires consistent flow rate of 5 - 10 volumes/24 hours, meter systems calibrated before study and checked twice daily during test period		
Aeration, if any	Not reported.			
Test vessel Material: (glass/stainless steel)	Glass beakers			
Size: Fill volume:	250 mL 200 mL	EPA requires: size 250 ml or 3.9 L fill 200 ml		

		Remarks		
Parameter	Details	Criteria		
Source of dilution water	The dilution water was freshwater obtained from an on-site laboratory well (40-m deep). The well water was			
	sand filtered, UV irradiated and aerated prior to use.	EPA requires soft reconstituted water or water from a natural source, not dechlorinated tap water.		
Water parameters: Hardness	136 mg/L as CaCO <sub>3</sub>	The hardness and pH were higher than recommended.		
pH	8.4-8.7	Results of the analysis of the well		
Dissolved oxygen	8.2-8.4 mg/L (≥93% saturation)	water on July 31, 2002 for pesticides, organics, and metals are provided in Appendix 3, pp. 25-26.		
Temperature	19.5-20.8°C	provided in Appendix 3, pp. 20 20.		
Total Organic Carbon	<1.0 mg/L	EPA requires: hardness: 40 - 48 mg/L as CaCO <sub>3</sub>		
Particulate matter	Not reported	pH: 7.2 - 7.6  -Temperature: 20°C (measured		
Metals	See Appendix 3, p. 26.	continuously or if water baths are used, every 6 hr, may not vary > 1°C		
Pesticides	<lod< td=""><td>Dissolved oxygen: Static: <math>\geq</math> 60% during 1<sup>st</sup> 24 hr and <math>\geq</math> 40% during 2<sup>nd</sup> 24 hr</td></lod<>	Dissolved oxygen: Static: $\geq$ 60% during 1 <sup>st</sup> 24 hr and $\geq$ 40% during 2 <sup>nd</sup> 24 hr		
Chlorine	Not reported	Flow-through: ≥60%		
Number of organisms per replicate Solvent control: Negative control:	N/A 20	The biomass loading rate was not specified.		
Treatments:	20	EPA requires 5 treatment levels plus control with a minimum of 20 daphnid per treatment. Biomass loading rate for static $\leq 0.8$ g/L at $\leq 17^{\circ}$ C, $\leq 0.5$ g/L at $\geq 17^{\circ}$ C; flow-through: $\leq 1$ g/L/day.		
Number of replicates Solvent control: Negative control: Treatments:	N/A 2 2			

		Remarks		
Parameter	Details	Criteria		
Treatment concentrations nominal:  measured:	0 (negative control), 0.31, 0.63, 1.3, 2.5, 5.0 and 10 ppm < 0.201 ( <loq, 0.30,="" 0.60,="" 1.2,="" 2.3,="" 4.7="" 9.5="" a.i.<="" and="" control),="" ppm="" td=""><td colspan="3">Mean-measured concentrations are provided in Table 1, p. 17.  Concentrations were stable during the 48-hour study.</td></loq,>	Mean-measured concentrations are provided in Table 1, p. 17.  Concentrations were stable during the 48-hour study.		
	<b>a.i.</b>	EPA requires a geometric series with each concentration being at least 60% of the next higher one.		
Solvent (type, percentage, if used)	N/A			
		EPA requires solvents not to exceed 0.5 mL/L for static tests or 0.1 mL/L for flow-though tests.		
Lighting	16 hours light/8 hours dark with a 30-minute transition period.	Light intensity was approximately 203 lux at test initiation (p. 13).  EPA requires 16 hours light, 8 hours dark.		
Feeding	Animals were not fed during testing.	EPA/OECD requires: No feeding during the study		
Stability of chemical in the test system	Verified. Recoveries were 91.9-96.8% of nominal concentrations at 0 hours and 90.2-98.6% of nominal at 48 hours (Table 1, p. 17).			
Recovery of chemical	96.0-102% of nominal	Based on quality control matrix		
Level of Quantitation	0.201 ppm a.i.	spikes fortified at 0.300, 1.50, or 10.0 ppm and analyzed concurrently with the samples		
Level of Detection	Not reported	(Appendix 4.5, p. 32).		
Positive control {if used, indicate the chemical and concentrations}	N/A			
Other parameters, if any	N/A			

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## 2. Observations:

Table 2: Observations

Criteria		Remarks  Criteria		
	Details			
Parameters measured including the sub-lethal effects	Mortality/immobility and sub- lethal effects			
Observation intervals	After 2, 24, and 48 hours			
Were raw data included?	Yes, sufficient			
Other observations, if any	N/A			

## II. RESULTS AND DISCUSSION

## A. MORTALITY

By 48 hours, mortality/immobility was 5% in the control group, 0% in the 0.30 through 2.3 ppm a.i. groups, 25% in the 4.7 ppm a.i. group, and 100% in the 9.5 ppm a.i. group (Table 4, p. 20). The 48-hour  $EC_{50}$  (with 95% C.I.) was 5.7 (4.7-9.5) ppm a.i. (Table 5, p. 21).

Table 3: Effects of Thidiazuron Technical on mortality/immobilization of Daphnia magna.

Treatment, ppm a.i. Measured and (nominal) concn.	No. of organisms	Observation period							
		2 ]	Hours	24 Hours		48 Hours			
		No.	%	No.	%	No.	%		
Negative Control	20	0	0	0	0	1	5		
0.30 (0.31)	20	0	0.	0	0	0	0		
0.60 (0.63)	20	0	0	0	0	0	0		
1.2 (1.3)	20	0	0	0	0	0	0		
2.3 (2.5)	20	0	0	0	0	0	0		
4.7 (5.0)	20	0	0	0	0	5	25		
9.5 (10.0)	20	0	0	16	80	20	100		
NOEC, ppm a.i.		Not reported		Not reported		Not reported			
LOEC, ppm a.i.		Not reported		Not reported		Not rep	Not reported		
LC/EC <sub>50</sub> (95% C.I.), ppm a.i.		Not reported		7.6 (4.7-9.5)		5.7 (4.7-9.5)			

# B. SUB-LETHAL TOXICITY ENDPOINTS:

No signs of toxicity were observed after 2 hours of exposure at any test level (Table 4, p. 20). After 24 hours, all surviving daphnids from the 9.5 ppm a.i. group were lethargic. After 48 hours, 1/20 surviving daphnids from the 2.3 ppm a.i. group and 6/15 surviving daphnids from the 4.7 ppm a.i. group were lethargic, and all daphnids from the 9.5 ppm a.i. group were either dead or immobile.

Table 4: Sub-lethal Effects of Thidiazuron Technical on Daphnia magna.

Treatment, ppm a.i. Measured and (nominal) concn.	Observation period						
	24	4 hours	48 hours				
	endpoint	% affected <sup>a</sup>	endpoint	% affected <sup>a</sup>			
Negative Control	Appear normal	0	Appear normal	0			
0.30 (0.31)	Appear normal	0	Appear normal	0			
0.60 (0.63)	Appear normal	0	Appear normal	0 .			
1.2 (1.3)	Appear normal	0	Appear normal	0			
2.3 (2.5)	Appear normal	0	Lethargic	5			
4.7 (5.0)	Appear normal	0	Lethargic	40			
9.5 (10.0)	Lethargic	100		_			
NOEC, ppm a.i.	Not determined		Not determined				
LOEC, ppm a.i.	Not determined		Not determined				
EC <sub>50</sub> (95% C.I.), ppm a.i.	Not determined		Not determined				

<sup>&</sup>lt;sup>a</sup> The percent of affected daphnia was reviewer-calculated from number affected based on number of surviving daphnids.

## C. REPORTED STATISTICS:

The  $LC_{50}/EC_{50}$  values for mortality/immobility (with 95% C.I.) were calculated using the binomial probability method (Stephan, C.E., 1978) and mean-measured concentrations (p. 14). The study authors did not report NOEC or LOEC values, or an  $EC_{50}$  value based on sub-lethal effects.

# 48-Hour

LC<sub>50</sub>/EC<sub>50</sub>: 5.7 ppm a.i. NOEC: Not determined LOEC: Not determined 95% C.I.: 4.7-9.5 ppm a.i.

<sup>- 100%</sup> Mortality/immobility

# D. VERIFICATION OF STATISTICAL RESULTS:

The 48-hour  $LC_{50}$  was determined for dead/immobile daphnids using the binomial probability method via TOXANAL statistical software, as this method provided a sound 95% confidence interval, compared to the probit and moving average methods. A NOEC was determined using Fisher's Exact Test via TOXSTAT statistical software based on mortality/immobility data. A NOEC was also visually determined as the highest concentration which exhibited no sub-lethal effects, the more sensitive endpoint. All toxicity values were determined in terms of the reported mean-measured treatment concentrations.

# 48-Hour

LC<sub>50</sub>/EC<sub>50</sub>: 5.7 ppm a.i. 95% C.I.: 4.7-9.5 ppm a.i.

NOEC: 1.2 ppm a.i. (based on sub-lethal effects)

LOEC: 2.3 ppm a.i.

Endpoints affected: Mortality/immobility and sub-lethal effects

Most sensitive endpoint: Sub-lethal effects

### E. STUDY DEFICIENCIES:

There were no significant deviations from U.S. EPA guideline §72-2 that affected the acceptability or validity of this study.

#### F. REVIEWER'S COMMENTS:

The reviewer's conclusions were identical to those reported by the study authors. However, the study authors did not report a NOEC or LOEC value based on mortality/immobility or sub-lethal effects, therefore the reviewer determined these values using appropriate statistical methods. Consequently, the reviewer determined NOEC and LOEC values reported in the Conclusion and Executive Summary sections of this DER are based on the reviewer's statistical verification and the reported sub-lethal effects data.

The test solutions appeared clear and colorless at test initiation and termination (p. 11).

## G. CONCLUSIONS:

This study is scientifically sound, fulfills U.S. EPA guideline §72-2, and is classified as CORE. Based on the results of this study, Thidiazuron Technical is categorized as moderately toxic to the Cladoceran, *Daphnia magna*, on an acute toxicity basis. The 48-hour NOEC and LOEC values were 1.2 and 2.3 ppm a.i., respectively, based on sub-lethal effects data, the most sensitive endpoint.

### 48-Hour

LC<sub>50</sub>/EC<sub>50</sub>: 5.7 ppm a.i. 95% C.I.: 4.7-9.5 ppm a.i.

NOEC: 1.2 ppm a.i. (based on sub-lethal effects)

LOEC: 2.3 ppm a.i.

Endpoints affected: Mortality/immobility and sub-lethal effects

Most sensitive endpoint: Sub-lethal effects

#### III. REFERENCES:

Organization for Economic Cooperation and Development. 1984. Guideline 202: Daphnia sp. Acute Immobilisation Test and Reproduction Test. OECD Guideline for Testing of Chemicals. Updated Guideline,

adopted April, 1984.

- U.S. Environmental Protection Agency. 1996. Aquatic Invertebrate Acute Toxicity Test, Freshwater Daphnids. Series 850 Ecological Effects Test Guidelines (draft), OPPTS Number 850.1010.
- ASTM Standard E729-88a. 1994. Standard Guide for Conducting Acute Toxicity Tests with Fishes, Macroinvertebrates, and Amphibians. American Society for Testing and Materials.
- APHA, AWWA, WPCF. 1998. Standard Methods for the Examination of Water and Wastewater. 20th Edition, American Public Health Association. American Water Works Association. Water Pollution Control Federation, New York.
- Stephan, C.E. 1978. U.S. EPA. Environmental Research Laboratory, Duluth, Minnesota. Personal communication.
- Thompson, W.R. 1947. Bacteriological Reviews. Vol. II, No. 2. Pp. 115-145.
- Finney, D.J. 1971. Statistical Methods in Biological Assay. Second edition. Griffin Press, London.
- Stephan, C.E. 1977. Methods for Calculating and LC50, Aquatic Toxicology and Hazard Evaluations. American Society for Testing and Materials. Publication Number STP 634, pp 65-84.

# APPENDIX 1. OUTPUT OF REVIEWER'S STATISTICAL VERIFICATION:

TOXANAL RESULTS: Calculated using the reported mean measured concentrations (Table 3, p. 19).

### LC50:

9.5	20	20	100	9.536742E-05
4.7	20	5	25	2.069473
2.3	20	0	0	9.536742E-05
1.2	20	0	0	9.536742E-05
. 6	20	0	0	9.536742E-05
. 3	20	0	0	9.536742E-05

THE BINOMIAL TEST SHOWS THAT 4.7 AND 9.5 CAN BE USED AS STATISTICALLY SOUND CONSERVATIVE 95 PERCENT CONFIDENCE LIMITS, BECAUSE THE ACTUAL CONFIDENCE LEVEL ASSOCIATED WITH THESE LIMITS IS GREATER THAN 95 PERCENT.

AN APPROXIMATE LC50 FOR THIS SET OF DATA IS 5.679563

WHEN THERE ARE LESS THAN TWO CONCENTRATIONS AT WHICH THE PERCENT DEAD IS BETWEEN 0 AND 100, NEITHER THE MOVING AVERAGE NOR THE PROBIT METHOD CAN GIVE ANY STATISTICALLY SOUND RESULTS.

 $\begin{tabular}{ll} \textbf{TOXSTAT RESULTS:} & \texttt{NOEC Determination based on mortality/immobilization data via } \\ \textbf{Fisher's Exact Test.} \\ \end{tabular}$ 

# SUMMARY OF FISHERS EXACT TESTS

GROUP IDENTIFIC		IFICA	TION	NUMBER EXPOSED			NUMBER DEAD	SIG (P=.05)	
			CONTROL		20	_	1		
1			0.3		20		0		
2			0.6		20		0		
3			1.2	3	20		0		23
4			2.3		20		0		
5			4.7		20		5		
6			9.5		20		20	*	