US ERA ARCHIVE DOCUMENT

DATA EVALUATION RECORD HONEY BEE - ACUTE ORAL LD₅₀TEST No OPP Guideline Applicable - Acute Oral

1. <u>CHEMICAL</u>: Clothianidin (TI-435) <u>PC Code No.</u>: 044309

2. TEST MATERIAL: TI-435 Metabolite MNG Purity: 99.2%

3. CITATION:

Author: Wilkins, P.

Title: TI-435 Metabolite MNG: Acute Oral Toxicity to Honey Bees

(Apis mellifera)

Study Completion Date: January 27, 2000

<u>Laboratory</u>: National Bee Unit

Central Science Laboratory

Sand Hutton, York Y041 1LZ, England

Sponsor: Takeda Chemical Industries Ltd

Development Department, Agro Company 13-10 Nihonbashi 2-chome, Chuo-Ku

Tokyo 103-8668, Japan

<u>Laboratory Report ID</u>: GQ3203(Study Number); 110056(Report Number)

DP Barcode: D278110

MRID No.: 45422428

4. Secondary Reviewer: Gabe Patrick, Biologist, EPA/OPPTS/OPP/EFED/ERB5

Signature: Halve Patrick Date: 3/5/03

Secondary Reviewer: Mike Rexrode, Ph.D., Senior Scientist, EPA/OPPTS/OPP/EFED/ERB5

Signature: M. Rexxoole Date: 3/5/03

5. <u>Secondary Reviewer:</u> Valerie Hodge, MSc, Senior Evaluation Officer

Environmental Assessment Division, PMRA

Signature: Valence Hodge Date: 3/20/03

2006172

6. STUDY PARAMETERS:

Scientific Name of Test Organism: Apis mellifera

Age or Size of Test Organism at Test Initiation: Adult worker bees, age not specified.

Type of Concentrations: Nominal

Definitive Study Duration: 48 hours

7. CONCLUSIONS:

The honey bee, *Apis mellifera*, was exposed to MNG, a metabolite of Clothianidin (TI-435), for 48 hours. There was 0% mortality in the control and treatment groups. There were no sub-lethal effects observed at any observation period for any bees exposed to MNG at 48 hours. The LD₅₀ was >153 μ g a.i./bee (highest dose tested) to the honey bee, *Apis mellifera*, on an acute oral toxicity basis. The NOAEL \geq 153 μ g a.i./bee. These toxicity values are based on analyzed content of MNG in the formulation [i.e., values were corrected for the actual purity of the test substance (99.2% ai) used in this study].

This study is scientifically sound and is classified as **Supplemental** for a non-guideline study.

Results Synopsis:

Results - Oral Test (48 h):

LD₅₀: >153 μg a.i./bee (highest dose tested)

NOAEL: ≥153 µg a.i./bee

95% C.I.: N/A

Probit Slope: N/A

8. ADEQUACY OF THE STUDY:

A. Classification: Supplemental

B. Rationale: This acute oral study is scientifically sound and is classified as

Supplemental because the study is a non-guideline study and does not

fulfill an OPP guideline requirement.

C. Repairability: NA

9. GUIDELINE (as compared to 141-1 or 850.3020) DEVIATIONS:

None noted.

10. SUBMISSION PURPOSE: This study was submitted to provide data on the acute oral toxicity of MNG, a metabolite of TI-435 for the purpose of chemical registration.

11. MATERIALS AND METHODS:

A. Test Organisms

Criteria	Reported Information		
Species: Species of concern (Apis mellifera)	Apis mellifera		
Age at beginning of test:	Adult worker bees, age not specified.		
Supplier:	Central Science Laboratory's National Bee Unit, York, England		
All bees from the same source?	Yes, all bees from a single colony.		

B. Test System

Criteria.	Reported Information		
Cage size adequate?	Cylindrical mesh cages, no size given.		
Lighting:	Darkness, except during observation periods.		
Temperature:	25 ± 1°C		
Relative humidity:	65 ± 5%		

C. Test Design

Criteria	Reported Information
Range finding test?	Bees were exposed to MNG at nominal concentrations of 0.109, 1.09, 10.9, and 109 µg ai/bee, with mortality rates of 3, 3, 0, and 0%, respectively. The definitive nominal concentrations were based upon these results.

Criteria	Reported Information		
Reference toxicant test?	Dimethoate, at concentrations of 0.063, 0.125, 0.25, and 0.5 µg ai/bee; 3 replicates with 10 bees/replicate.		
Method of administration:	 - Mixed with diet (50% w/v aqueous sucrose) - Doses offered to bees within 2 h after preparation - 200 μL/treatment group offered in glass feeder 		
Nominal doses:	0.181, 1.81, 18.1, and 181 μg a.i./bee		
Controls: Negative control and/or diluent/solvent control	Negative control.		
Number of colonies per group:	3 replicates per treatment group, 10 bees per replicate.		
Solvent: The following solvents: acetone, dimethylformamide, triethylene glycol, methanol, ethanol.	50% w/v aqueous sucrose		
Feeding:	Bees were starved for 1.5 to 2 hours prior to test initiation. After first four hour observations, the food solution containing MNG was removed and fresh aqueous sucrose (50% w/v) was provided ad libitum.		
Observations period:	Mortality and behavior observations: 4 hours after test was initiated, then 24 and 48 hours after treated diet was removed from cages.		

12. <u>REPORTED RESULTS:</u>

Guideline Criteria	Reported Information
Quality assurance and GLP compliance statements were included in the report?	Yes
Control performance:	0% mortality by 48 hours.
Raw data included:	Yes
Signs of toxicity (if any) were described?	Knocked down (alive but immobile) bees were observed early on in the range-finding test. This sub-lethal effects disappeared by 24 hours. No signs of toxicity were present in the definitive test.

Mortality - Oral Test

			Cumulative Number of Dead		
Dosage ^a	No. of		Hour of Study		
(μg ai/bee)	bees	Rep.	4	24	48
Negative Control	10. 10. 10.	1. 2. 3.	0. 0. 0.	0. 0. 0.	0. 0. 0.
(0.181) 0.18 0.18 0.18	10. 10. 10.	1. 2. 3.	0. 0. 0.	0. 0. 0.	0. 0. 0.
(1.81) 1.8 1.8 1.6	10. 10. 10.	1. 2. 3.	0. 0. 0.	0. 0. 0.	0. 0. 0.
(18.1) 18 18 18	10. 10. 10.	1. 2. 3.	0. 0. 0.	0. 0. 0.	0. 0. 0.

	No. of	Rep.	Cumulative Number of Dead Hour of Study		
Dosage* (µg ai/bee)					
	bees		4	24	48
(181)					
171	10.	1.	0.	0.	0.
158	10.	2.	0.	0.	0.
131	10.	3.	0.	0.	0.

The second secon		Trans.	Cumulative Number of Dead Hour of Study		
Dosage ^a	No. of				
(μg ai/bee)	bees	Rep.	4	24	48
Reference Toxican	t (Dimethoa	te)			
Negative Control	10. 10. 10.	1. 2. 3.	0. 0. 0.	0. 0. 0.	0. 0. 0.
(0.063) 0.061 0.063 0.057	10. 10. 10.	1. 2. 3.	0. 0. 0.	2. 1. 1.	2. 1. 1.
(0.125) 0.12 0.11 0.12	10. 10. 10.	1. 2. 3.	0. 0. 0.	2. 1. 4.	2. 1. 4.
(0.25) 0.23 0.19 0.21	10. 10. 10.	1. 2. 3.	0. 0. 0.	10. 3. 6.	10. 3. 6.
(0.50) 0.43 0.35 0.41	10. 10. 10.	1. 2. 3.	0. 0. 0.	10. 9. 8.	10. 10. 9.

^a Nominal dosages are listed in parentheses. Nominally consumed dosages, based on percent diet consumed, are listed per replicate.

Note: There was a glitch in the above table whereby data numbers appear on a single line instead of separate lines. To prevent this I put a "." after each number.

<u>Observations</u>: There was no mortality observed in the control or the MNG treatment groups. No bees were observed to be knocked down or stumbling at any of the observation times in the main oral dosing test by 48 hours.

Percent mortality in the Dimethoate nominal 0.063, 0.125, 0.25, and 0.50 μg a.i./bee groups was 13, 23, 63 and 97%, respectively, by 48 hours.

Statistical method: The MNG LD₅₀ value and the NOEL value were determined visually based on mortality data, since no treatment group exceeded 50% mortality. The study author determined these values based on the nominally consumed dosage concentrations.

Reported Statistical Results

LD₅₀: >153 μ g a.i./bee NOEL: \geq 153 μ g a.i./bee 95% C.I.: N/A

Probit Slope: N/A

13. <u>VERIFICATION OF STATISTICAL RESULTS</u>:

Method: The mortality data were visually inspected to determine the NOAEL and LD₅₀.

Verified Results

 LD_{50} : >153 µg a.i./bee

95% C.I.: N/A

NOAEL: ≥153 µg a.i./bee

Probit Slope: N/A

14. REVIEWER'S COMMENTS:

The reviewer's conclusions were identical to the study author's. The MNG LD₅₀ was >153 µg a.i./bee to the honey bee, *Apis mellifera*, on an acute oral toxicity basis. The NOAEL \geq 153 µg a.i./bee.

The nominally consumed test concentration refers to the nominal dose consumed by the bees. The dose consumed was determined by the weight of dose remaining in the glass feeders after the 4 h treatment period and comparing this weight to the weight of the known volume of the test solutions at the beginning of the test. This is not a measured (analyzed) dose per se.

The toxicities values provided in this study by the author and reviewer were corrected for the actual purity of the test substance (99.2% ai) used in this study.

There was no data provided on the stability of the test substance in solution for this study. The test substance was assumed to be stable by the author and reviewer.

MNG (a metabolite of clothianidin) was classified as "Virtually non-toxic" to honey bees on an acute oral basis by the author according to the following ICBB (1985) categorization:

 $> 100 \mu g a.i./bee$

Virtually non-toxic

10-100 μg a.i./bee 1-10 μg a.i./bee Slightly toxic Moderately toxic

< 1.0 µg a.i./bee

Highly toxic

OPP does not have a categorization scheme for acute oral toxicity to honey bees.

This study was conducted in accordance with UK Good Laboratory Practice Regulations, USEPA Title 40 CFR 160, Japan Ministry of Agriculture, Forestry and Fisheries, and OECD Principles of Good Laboratory Practices. OPP does not have or require an acute oral LD₅₀ study for honey bees.

15. REFERENCES:

1985. International Commission for Bee Botany Third Symposium on the "Harmonization of methods for testing the toxicity of pesticides to bees".

1992. European and Mediterranean Plant Protection Organization. "Guideline on test methods for evaluating the side-effects of plant protection products on honey bees" <u>EPPO Bulletin 22</u>, 203-215

1996. Ministry of Agriculture, Fisheries and Food (UK), Pesticides Safety Directorate and the Health and Safety Executive, "The Registration Handbook Volumes 1 and 2, Pesticides, Biocides, Plant Protection Products, A guide to the policies, procedures and data requirements relating to their control within the United Kingdom".

1997. OECD (Draft April-Adopted 21/09/98). "OECD Guidelines for the testing of chemicals. Proposal for a new guideline 213. Honeybees, acute oral toxicity test."

EAD Assessment of USEPA DER

Reviewer: Valerie Hodge

Date: November 7, 2002

PMRA Submission Number: 2001-1293

Study Type: TI-435 Metabolite MNG: Acute Oral LD₅₀ - Honeybee [*Apis mellifera*], PMRA DATA CODE 9.2.4.2, EPA MRID Number 45422428, OECD Data Point IIA 8.7.1, EPA Guideline - none.

Reviewing Agency: U.S. EPA

EAD Summary:

The honey bee, *Apis mellifera*, was exposed to MNG, a metabolite of clothianidin (TI-435), for 48 hours (4 hour exposure followed by observation to 48 hours) at mean nominal doses of 0.18, 1.7, 18, and 153 µg a.i./bee. There was 0% mortality in the control and treatment groups. There were no sub-lethal effects observed at any observation period for any bees exposed to MNG. The acute oral LD₅₀ was >153 µg a.i./bee (highest dose tested) to the honey bee, *Apis mellifera*. The NOEL was \geq 153 µg a.i./bee (based on mortality and sublethal effects). These toxicity values were corrected for the actual purity of the test substance (99.2% ai) used in this study. MNG is "virtually non-toxic" to bees by ingestion.

Material and Methods:

Doses of MNG were prepared in 50% w/v aqueous sucrose. Adult worker bees (*Apis mellifera* L.), 3 replicates of 10 per treatment and control, were exposed in mesh cages to a given concentration (or control) of MNG for 4 hours (200 μ L per 10 bees in a glass feeder). After 4 hours, clean feeding solution replaced treatment solutions for the remainder of the study period. Nominal doses were 0.181, 1.81, 18.1, and 181 μ g a.i./bee. Actual nominal doses were determined by weighing the feeding solution before and after exposure to bees, and determining a mean dose per bee. Doses were also corrected for the purity of the test substance (99.2% ai). Actual mean nominal doses were 0.18, 1.7, 18, and 153 μ g a.i./bee. Observations for mortality and behavior (knockdown or stumbling) were made at 4 hours after test was initiated, then 24 and 48 hours after treated diet was removed from cages.

Bees were also exposed to dimethoate, as a reference toxicant, at nominal doses of 0.063, 0.125, 0.25, and $0.50 \mu g$ a.i./bee.

Results:

There was 0% mortality in the control and treatment groups at all observation times. There were no sub-lethal effects observed at any observation period for any bees exposed to MNG.

Statistical Results

LD₅₀: >153 μg a.i./bee NOEL: ≥153 µg a.i./bee 95% C.I.: N/A Probit Slope: N/A

Verified statistical results agreed with reported results.

Results for dimethoate toxicity were consistent with previous studies. Percent mean mortality in the dimethoate (actual) nominal 0.06, 0.12, 0.21, and 0.40 µg a.i./bee groups was 13, 23, 63 and 97%, respectively, by 48 hours (48 hour LD₅₀, 0.16 μ g dimethoate/bee).

EAD comments:

The EAD evaluator agrees with the conclusions reached by the U.S. EPA evaluator.

EAD Conclusion:

Based on the results of this study, and the criteria of the International Commission for Bee Botany (1985), MNG is "virtually non-toxic" to bees by ingestion.

Reference: 1985. International Commission for Bee Botany Third Symposium on the "Harmonization of methods for testing the toxicity of pesticides to bees".

Signatures:

Primary Reviewer:

Valerie Hodge

Date: November 7, 2002

Secondary Reviewer: Hemendra Mulye

Date: