

US EPA ARCHIVE DOCUMENT



271DD

009109

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
WASHINGTON, D.C. 20460

OFFICE OF
PESTICIDES AND TOXIC SUBSTANCES

MEMORANDUM

SUBJECT: EPA Reg. No./File Symbol 279-GNIR
CYNOFF 2 EC Insecticide

FROM: William S. Woodrow WSW 5-17-89
Precautionary Review Section E 5/17/89
Registration Support Branch
Registration Division (H75-G5C)

TO: George LaRocca (PM 15)
Insecticide-Rodenticide Branch
Registration Division (TS-767C)

APPLICANT: FMC Corp.
Agricultural Chemical Group
2000 Market St.
Philadelphia, PA 19103

FORMULATION FROM LABEL:

Active Ingredient(s):	% by wt.
<u>α-(+)-cyano-(3-phenoxyphenyl)methyl(+)-cis,</u>	
<u>trans-3-(2,2-dichloroethyl)-2,2-dimethyl-</u>	
<u>cyclopropane carboxylate</u>	<u>24.8</u>
<u>Inert Ingredient(s):</u>	<u>75.2</u>
Total	100.0%

BEST AVAILABLE COPY

157

BACKGROUND:

The FMC Corp. submitted Acute oral, dermal, Primary eye and dermal irritation, and Dermal sensitization studies, to support Registration of CYNOFF 2 EC Insecticide. MRID NOS. USED WERE: 406809-02 through 406809-06.

RECOMMENDATION:

1) The 5 acute studies submitted by FMC are acceptable to RSB/PRS. The Dermal sensitization study was designated Core minimum data, due to the lack of a positive control experiment.

2) An Acute inhalation toxicity evaluation of CYNOFF 2 EC Insecticide, will be necessary.

LABELLING:
NOTE: Upon submission of an acceptable acute inhalation study, Precautionary labeling may require revision.

LABELLING:

- 1) The CAUTION signal word is appropriate.
- 2) The Precautionary Statements are acceptable.
- 3) The Statements of Practical Treatment are acceptable.

DATA REVIEW FOR ACUTE ORAL TOXICITY TESTING (§81-1)

Product Manager: (15) Reviewer: Woodrow M. Waller
 MRID No.: 406809-02 Report Date: 5-16-89
 Testing Facility: FMC Tox. Lab. Report No. A87-2596
 Author(s): C. Freeman
 Species: Rat, Sprague Dawley
 Age: young adult Observation Days (Post Exposure): (14); other ()
 Weight: M 211-248, F 201-237g
 Source: Taconic Farms, N.Y.
 Test Material: Cypermethrin (Cisnelt) 2E
 Quality Assurance (40 CFR §160.12): satisfactory

Conclusion:

- LD₅₀ (mg/kg): Males = 1085 (645-1524); Females = 1246 (1048-1443); Combined = 1105 (841-1369)
- The estimated LD₅₀ is 1105 (841-1369)
- Tox. Category: III. Classification: Guidelines

Procedure (Deviations From §81-1): 25% (w/v) sol. in tap water, by gavage to groups of 5M & 5F rats. Animals observed for mortality & toxicity, to 14 days or longer if exhibiting clinical signs. Body wts. Necropsies on dying/dead, and surviving animals.

Reported Mortality

M	DOSAGE (mg/kg)	F	(NUMBER KILLED/NUMBER TESTED)		
			Males	Females	Combined
	2000	2000	4/5	5/5	9/10
	1000	1500	4/5	4/5	9/10
	950	1300	3/5	5/5	7/10
	900	1225	1/5	3/5	4/10
	800	1200	0/5	1/5	1/10
	500	1000	1/5	0/5	1/10

Symptomology & Gross Necropsy Findings:

Clinical signs included tremors, hypersensitivity to touch, ataxia, clonic convulsions, abdominal joint staining & oral discharge. All survivors gained weight by termination. The only gross intestinal necropsy finding was blood in the intestine of one rat in the 1200 mg/kg group.

BEST AVAILABLE COPY

DATA REVIEW FOR ACUTE DERMAL TOXICITY TESTING (§81-2)

Product Manager: (15)
 MRID No.: 406809-03
 Testing Laboratory: Emc Tox. Lab.
 Author(s): C. Freeman
 Species: Rabbits, M & F, White
 Sex: 5M & 5F Wt.: M 2.32-2.70, F 2.46-2.81 kg.
 Test Material: Cypermethrin (Cyvalif) 2 EC
 Quality Assurance (40 CFR §160.12): Satisfactory

Reviewer: Woodrow Weller
 Report Date: 5-17-79
 Report No. A97-2548

Summary:

- LD50 (mg/kg): Males = —; Females = —; Combined = —;
- The estimated LD50 is > 2.00 mg/kg.
- Tox. Category: III. Classification: Guidelines

Procedure (Deviations from §81-2): 2000 mg/kg to clipped backs of 5M & 5F rabbits; 4x4" gauze pad/tape. Occluded & taped. 24 hr contact. Sites washed, animals observed for mortality & irritation at 0.5, 1, 2, 3, 4, 6 hrs days 1 & 2, 2x daily to 14 days & necropsy

Results:

Reported Mortality

DOSAGE (mg/kg)	(NUMBER KILLED/NUMBER TESTED)		
	Males	Females	Combined
<u>2000 mg/kg</u>	<u>0/5</u>	<u>0/5</u>	<u>0/10</u>

Symptomology & Gross Necropsy Findings:

During 1st 3 days: loss of muscle control, decreased locomotion, recumbency, hyperaesthetic to touch, thrashing in cage, belching, and diarrhea. One sacrificed female (mechanical injury) showed brown material in stomach. Remaining animals normal.

BEST AVAILABLE COPY

A

003109

DATA REVIEW FOR ACUTE EYE IRRITATION TESTING (§81-4)

Product Manager: (15)
 MRID No.: 406909-04
 Testing Laboratory: Env. Tok. Lab.
 Author(s): C. Freeman
 Species: Rabbit, NZ White
 Sex: Female Weight: 2.36 - 3.51 kg.
 Source: Hudson Co.
 Dosage: 0.1ml
 Test Material: Supernethin (Synalt) 2 EC 27% a.i.
 Quality Assurance (40 CFR §160.12): Satisfactory

Reviewer: Woodrow
 Report Date: 5-17-79
 Report No. A27-2549

Summary:

Tox. Category: III Classification: Guidelines

Procedure (Deviation from ~~method~~): 0.1ml instilled into eye of 9 rabbits.
6 treated eyes unswathed, 3 eyes swathed 2 100ml tap water, off test.
20-30 sec after treatment. Eyes were examined and record according to
Diagn. at 1, 2, 4, 48 & 72 hrs.

Results:

	Observations (number "positive"/number tested)							
	Hour	Days						
	1	1	2	3	4	7	14	21
Cornea Opacity	0/9	0/9	0/9	0/9				
Iris	0/9	0/9	0/9	0/9				
Conjunctivae Redness	6/9	7/9	3/9	0/9				
Chemosis	0/9	1/9	0/9	0/9				
Discharge	7/9	2/9	1/9	0/9				

Comments: No iris or corneal involvement. All irritations
absent by day 3.

BEST AVAILABLE COPY

5

DATA REVIEW FOR SKIN IRRITATION TESTING (101-5)

Product Manager: (15)
 Reviewer: Woodrow
 HRID No.: 406909-05
 Report Date: 5-17-79
 Testing Laboratory: FMC 70A Lab.
 Report No.: A77-2545
 Author(s): C. Freeman
 Species: Rabbit, Al 2 white
 Age: Young adult
 Sex: 3M 3F
 Weight: 2.72 - 2.82 kg
 Dosage:
 Test Material: Cypermethrin (Cythoff) 2 EC 2770 AE
 Quality Assurance (40 CFR 160.121): Satisfactory

Summary:

The Primary Irritation Index = P.I. Index = 0.40
 Toxicity Category: IV
 Classification: Guidelines

Procedure (~~Deviation from 101-5~~): 0.5 ml to clipped back of each of 6 rabbits, under 2" sq. gauze - Entire back wrapped in semi-occlusive chesulth bandage - 4 hr exposure - Dressing removed, sites wiped. Scoring of sites beginning 30 min after unwrapping, and at 24, 48, 72 hrs, and on days 4 & 5, according to Draeg.
 Results:

Scoring interval	Irritation score
4.5 hrs	0.2
24.0 hrs	0.3
48.0 hrs	0.5
72.0 hrs	0.4
day 4	0.3
day 5	0.0

P.I. Index = 0.4; practically not an irritant

Special Comments:

BEST AVAILABLE COPY

6

DATA REVIEW FOR SKIN SENSITIZATION TESTING (§81-6)

Product Manager: (15)
 MRID No.: 406900-06
 Testing Laboratory: EMC Lab. Lab.
 Author(s): C. Freeman
 Species: guinea pigs, Hartley
 Sex: not stated
 Source: Harlan Rac. Animals
 Test Material: Cypermethrin (Cyvalb) 2 EC 379, 6E.
 Positive Control Material: none
 Quality Assurance (40 CFR §160.12): Laboratory
 Reviewer: Woodrow Miller
 Report Date: 5-17-89
 Report No. A87-2550
 Method: Modified Buchler.

Summary:

1. This product is / (Is not a dermal sensitizer.)
2. Classification: Case minimum data (one treated ctrl.)

Procedure (Deviation from §81-6):

Induction - 0.3 ml to each of 20 Hill Top Chamber. Chamber applied to clipped dorsal sites of 20 g.p./secured a tape. Tape was chamber. 6 hr exposure - sites affixed, wiped. ~~none~~ G.p. dorsal then, back/punch, to total 3 applications - 14 day rest period, all test animals challenged on a single site on the right shoulder. 10 additional animals (challenge only), each received 0.3 ml, for comparison - all patches removed later.

Test sites scored for irritation at 24 & 48 hrs, according to Dosing (system/dose):

0 = insignificant

2 = significant

1 = equivocal

BEST AVAILABLE COPY

One index for incidence, and one for severity.

Results: All animals remained healthy.

The only scoring for irritation, was 1/20 animals, for the test material challenge, at 24 hours; a single incidence (calculated to be 0.05), which was for a score of 1.0 No other irritation responses.

Conclusion: Cypermethrin 2 EC, did not irritate guinea pigs.