

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

004415

MEMORANDUM

OFFICE OF
PESTICIDES AND TOXIC SUBSTANCES

SUBJECT:

EPA Reg. No.: 279-2862; Carbofuran; miscellaneous

data; five acute studies

Caswell #: 160A Accession #: 256481

TO:

Jay Ellenberger

Product Manager (12)

Registration Division (TS-767)

THRU:

12 4/sx/se Robert P. Zendzian Ph.D.

Acting Head, Review Section IV

Toxicology Branch

Hazard Evalualtion Division (TS-769)

FROM:

William Dykstra, Ph.D.

Toxicology Branch

Hazard Evaluation Division (TS-769)

William Dykatras 18-769) 4124/85 Mijn(6-16)

Requested Action:

Review acute oral LD50 studies on carbamate and phenolic metabolites of carbofuran to determine relative toxicity.

Conclusions:

The acute oral ${\rm LD}_{50}$ studies are acceptable and support the registration of carbofuran.

Background:

The registrant, FMC, submitted these additional studies to Agriculture Canada in order that a determination of the toxicological significance of carbamate and phenolic metabolites could be made.

Review:

1. Acute oral LD_{50} of FMC 18209 (3-hydroxy carbofuran) technical in rats (FMC # A83-1136; 4/12/84).

Groups of 10 male and 10 female, Sprague-Dawley rats were orally gavaged with graded doses of test material in corn oil.

Observation was 14 days.

Results:

Mortality data are shown below:

Male		Female	
Dosage mg/kg	% mortality	Dosage mg/kg	% mortality
25 20 18	80 30 10	18 15 10 8 7.5 7.0 5.0	100 70 60 40 55 45

Male $LD_{50} = 21.9 \text{ mg/kg} (19.9 - 23.8)$

Female $LD_{50} = 8.3 \text{ mg/kg } (6.8 - 9.9)$

Toxic Signs: Tremors, oral discharge, stained fur.

Body Weight: All but two surviving animals gained weight.

Necropsy: No gross lesions except for blood in intestine of one rat.

Toxicity Category I: Danger.

Classification: Core minimum data. 4

2. Acute oral LD₅₀ study with FMC 1781 Technical (3-Keto carbofuran) in rats (FMC study #: A83-1137; 4/12/84).

Groups of 10 male and female Sprague-Dawley rats were orally gavaged with graded doses of test material in corn oil.

Observations were for 14 days.

Results: Mortality data are shown below:

Male			Female	
Dosage mg/kg	% mortality	Dosage mg/kg	% mortality	
200 120	100 80	150 120	100 80	
110 100	90 20 .	100 70	40 30 <i>Z</i>	
70	10			

Male $LD_{50} = 108 \text{ mg/kg } (94.5-121)$

Female $LD_{50} = 93.1 \text{ mg/kg} (76.2-110)$

Combined $LD_{50} = 107 \text{ mg/kg (91.1-123)}$

Toxic Signs: Tremors, oral discharge, stained fur and

decreased locomotion.

Body Weight: Survivors gained weight.

Necropsy: No gross lesions.

Toxicity Category II: Warning.

Classification: Core minimum data.

3. Acute oral LD50 study with FMC 1649% technical (3-hydroxy-7-phenol) in rats (FMC A83-1134; 3/29/84).

Groups of 10 males and 10 females Sprague Dawley rats received oral gavage graded doses of test material in corn oil.

Observation was for 14 days.

Results:

Mortality data are shown below:

Male		Female		
Dosage mg/kg	9	mortality	Dosage mg/kg	% mortality
2400 2000 1400		80 60 10	2000 1700 1400	90 50 20
Male	LD ₅₀	= 1916 mg/kg	(1656-2175)	
Female	LD ₅₀	= 1654 mg/kg	(1500-1807)	

Toxic signs: Prostation, decreased locomotion, nasal, ocular

and oral discharge.

Body weight: Survivors gained weight.

Necropsy: Blood in intestines of decedents.

Toxicity Category III: Caution.

Classification: Core minimum data.

4. Acute oral LD₅₀ study with FMC 16490 technical (3-keto-7-phenol) in rats (FMC # A83-1135; 3/28/84).

Groups of 10 males and 10 females Sprague Dawley rats were orally gavaged with graded dosages of test material in corn oil. Observations was for 14 days.

Results:

No deaths at 300 and 800 mg/kg of test material.

 $LD_{50} > 800 \text{ mg/kg (both sexes)}$

Toxic Signs: Lacrimation and decreased locomotion in the 800 mg/kg.

Body Weight: All rats gained weight.

Necropsy: No gross lesions.

Toxicity Category III: Caution.

Classification: Core minimum data.

5. Acute oral LD₅₀ study with FMC 10272 technical (7-phenol) in rats (FMC # A83-1133; 3/28/84).

Groups of 10 males and 10 females Sprague-Dawley rats were orally gavaged with graded dosages of test material undiluted. Observations was for 14 days.

Results:

Mortality data are shown below:

Male		Female	
Dosage mg/kg	% mortality	Dosage mg/kg	% mortality
3000	80	2300	70
2300	40	1800	50
1800	10	1400	40
		1000	10

Male $LD_{50} = 2450 \text{ mg/kg} (2137-2764)$

Female $LD_{50} = 1743 \text{ mg/kg} (1362-2124)$

Toxic Signs: Prostation, tremors, discharge, decreased
locomotion.

Body Weight: Survivors gained weight.

Necropsy: Blood in intestine of decedents.

Toxicity Category III: Caution.

Classification: Core minimum data.