



Classification: Core-Minimum Data

- (1) although only 1 dose level was tested, the results are adequate since it is apparent that the maximum concentration was generated (the test material adhered to the chamber walls)

2. Acute Eye Application - (Hazleton Laboratories, Report Date 5/13/58, submitted by Diamond Alkali Co in PP#411)

3.0 mg of (presumably) 100% technical was placed into the conjunctival sac of the left eye of each of 3 rabbits. The treated eye was held closed for  $\approx$  30 sec., after which an immediate reading was made. Observations for gross signs of eye irritation were continued at 1, 4 and 24 hours, and daily thereafter for an additional 6 days.

Results

Immediately following application there was a mild degree of irritation which was characterized by erythema and vascularization of the sclera and nictitating membrane, accompanied by lacrimation. Irritation persisted through the 4 hr interval. The eyes appeared grossly normal after 24 hours.

TOX Category: III

Classification: Supplementary Data

- (1) an inadequate amount of the test material was instilled into the eye.

The following studies were submitted in PP#411 and reviewed by G.E. Whitmore on 11/17/64. The results of the studies and their classification are listed below.

1. Acute Oral Administration - (Hazleton Laboratories, Report Date 5/13/58, submitted by Diamond Alkali in PP#411)

LD<sub>50</sub> > 3160 mg/kg (no mortalities occurred)

TOXIC Signs: depression

Necropsy: unremarkable

TOX Category: III

Classification: Core-Minimum Data

- (1) although the observation period was only 7 days, the results adequately reflect the acute oral toxicity when one considers the results of other toxicity studies.

2. Acute Dermal Application - (Hazleton Laboratories, Report Date 5/13/58, submitted by Diamond Alkali in PP#411)

LD<sub>50</sub> > 10000 mg/kg (no mortalities occurred)

Toxic Signs: mild to moderate erythema, slight edema: the irritation observed decreased in intensity but was still present at 7 days.

003297

Necropsy: unremarkable

TOX Category: III

Classification: Core-Minimum Data

- (1) although the observation period was only 7 days, the results adequately reflect the acute dermal toxicity when one considers the results of other toxicity studies.

Typists: TH

NET CONTENTS

POUNDS  
003297

EPA Reg. No. 677-

LOGO

DIAMOND SHAMROCK

90% DIMETHYL T

Active Ingredient:		
Dimethyl tetrachloroterephthalate .....		90.0%
Inert Ingredients: .....		<u>10.0%</u>
	TOTAL .....	100.0%

KEEP OUT OF REACH OF CHILDREN

CAUTION

SEE SIDE PANEL FOR ADDITIONAL CAUTIONS

DIAMOND SHAMROCK CORPORATION • AGRICULTURAL CHEMICALS DIVISION

CLEVELAND, OHIO 44114

Left Panel

CAUTION

Harmful if swallowed.

Avoid contact with skin, eyes and clothing. In case of contact, flush eyes with plenty of water for at least 15 minutes.

Wash thoroughly after handling or using.

DO NOT contaminate water, food, feed, seed, fertilizers or other pesticides by storage or disposal.

DO NOT reuse empty container. Dispose of in an incinerator or landfill approved for pesticide containers, or bury in a safe place.

Stay away from smoke or fumes.

Consult Federal, state or local disposal authorities for approved alternative procedures such as limited open burning.

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Right Panel

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