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

SEP 8 1987 OFFICE OF
PESTICIDES AND TOXIC SUBSTANCES

MEMORANDUM

SUBJECT: BLOCKADE cat and dog sprays

TO:

Mr. George LaRocca, PM 15

Registration Division (TS-767C)

FROM:

Byron T. Backus, Toxicologist ! by

Toxicology Branch (TS-769C)

THROUGH:

Section Head, Review Section III W. Law Quent

Toxicology Branch (TS-769C)

and

Theodore M. Farber, Ph.D., D.A.B.T.

Branch Chief

Toxicology Branch (TS-769C)

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EPA Record Nos. 197405, 197776

Project No. 7-0976

Tox. Chem. 346, 77A

EPA Reg. Nos. 2596-114 Hartz Blockade for Cats 2596-115 Hartz Blockade for Dogs

Action Requested:

The Registration Division has requested an expedited review of 6(a)(2) and toxicity data on a DEET + Fenvalerate formulation (Hartz Mountain Blockade) registered for spray-on use on cats There have been a considerable number of reports of adverse effects on these animals following label-use exposure, and part of the material received is a computerized listing from the National Animal Poison Control Center reporting on 162 calls received on this product between March 10 through July 17, 1987.

Because this memorandum has been expedited, a comprehensive review of the registrant's toxicity data has not yet been made, although the data have been examined and what appear to be some of the more relevant problems are indicated in this memorandum.

Background:

What appears to be the same formulation is registered under two EPA registration numbers, 2596-114 (use on cats) and 2596-115 (use on dogs). In both cases the label declaration for active ingredients gives N,N-Diethyl-m-toluamide as 8.55% with other isomers 0.45%, Cyano (3-phenoxyphenyl)methyl-4-chloro-alpha-(1-methylethyl)benzeneacetate (also known as pydrin and fenvalerate) as 0.09%. The remaining 90.91% of the formulation is listed as "inerts." Precautionary statements and label use directions (with the exception of which species is to be treated) are identical.

Comments and Recommendations:

1. The rat oral LD_{50} for N,N-diethyl-m-toluamide technical (90-100%) is reported (DEET Registration Standard) as 2.43 ml/kg for males and 1.78 ml/kg for females. Since the specific gravity of the technical is reported (Registration Standard) as 0.99-1.00, these values can also be expressed as 2.43 gm/kg and 1.78 gm/kg. If one ounce (28 grams) of this formulation is sprayed on a cat, this represents a total of 28 x 0.0855 = 2.39 gms of DEET. Assuming that cats are as susceptible to DEET as are rats (and in fact they may be even more so), the potential then exists for a cat - particularly a young animal weighing 1 kg or less to ingest an LD50 amount of this active by grooming. label directions for use specify spraying the product over the entire body (avoiding the eyes), including feet and tail. However, there is no indication that the Agency has received information as to what sort of dose (amount sprayed on the animal) would be involved. In theory, given the label directions for use, the contents of the entire can (7 oz) could be sprayed on a cat and this would not constitute misuse.

In a study conducted at Leberco Testing, Inc. the rat oral LD50 on the technical concentrate (14.286% DEET) of this Hartz product was reported as 6700 mg/kg, with 95% confidence limits of 5668 to 7919 mg/kg. However, this oral LD50 is based on the combined mortality for both sexes; it is noteworthy that all females receiving doses of 7 g/kg or more died, while only 4/15 males did so, suggesting females are more susceptible. Symptoms of toxicity included moderate to severe ataxia even at the lowest dose level (5 g/kg).

While the BLOCKADE formulation also contains 0.09% Pydrin, the available information indicates that this contributes very little to the toxicity of the formulation (the lowest oral LD50 value for technical Pydrin is approximately 75 mg/kg; for a product containing 0.09% as sole active this would extrapolate to approximately 10,000 times this value).

The IRB/TSS review of the reported incidents notes that "The most at-risk group would appear to be young kittens (especially female)."

- 2. The material submitted August 28, 1987 by the Hartz Mountain Corporation includes only two reports involving cats. Both were efficacy studies using the aerosol product; one with two treated animals and the other with only one. While no signs of adverse reactions were noted in either of these studies, the total number of animals involved (three) is clearly inadequate to demonstrate product safety for cats. Additionally, no information was provided as to the sexes or ages of these animals, or the dosage amounts of product applied.
- 3. While the labeling for the cat product makes a 21-day efficacy claim and states: "Under normal circumstances, repeat spray every 21 days" it is also stated that: "In areas or periods of heavy infestation it may be necessary to treat the animal more frequently." The implication is that a cat could be treated as often as necessary. This may, however, be a somewhat moot point as indications are that a single exposure to the product may be sufficient to kill a cat.
- 4. According to information on p. 123 of the ninth (1986) edition of Current Veterinary Therapy, DEET (in the form of DEET-containing products such as Off, Deep Woods Off) was associated with seven suspected cases of toxicosis in cats. These incidents apparently occurred as a result of treatment of pets with products normally used on humans. "In dogs and, especially, cats, primary signs included seizures, tremors, and sometimes emesis. Ataxia or lethargy occasionally occurred." Symptoms reported for cats by the National Animal Poison Control Center included increased salivation, tremors, vomiting, ataxia /incoordination and depression.

According to the Toxicology Branch one-liners, symptoms in rats following dosage with a product containing 22.95% DEET (also with dimethyl phthalate, 15.3%; Di-n-propyl isocinchomeronate, 1%; and N-octyl bicycloheptene dicarboximide 1%) included hypoactivity, ataxia, salivation, prostration and loss of righting reflexes. The oral LD50 for males was 4550 (3434-6029) mg/kg, and for females was 5450 (3759-7903) mg/kg.

5. After taking into account the data indicated above, as well as the considerable number of reported adverse incidences involving cats (109, with 5 deaths) following exposure to Blockade as reported to the National Animal Poison Control Center from March 10 through July 17, 1987, the Toxicology Branch is of the opinion that there is an insufficient safety factor associated with normal (following

label directions) use of EPA Reg. No. 2596-114 (Hartz Blockade for Cats).

- 6. The reported incidents involving dogs are also of considerable concern, although these appear to be less frequent and usually less severe than those involving cats. However, a considerable number of deficiencies are evident on examination of the label for 2596-115 (Hartz Blockade for Dogs). These include the following:
 - i) Sick and debilitated dogs are not excluded from treatment, as well as puppies and possibly toy breeds with large body surface to volume ratios (a death was reported in a 4 lb 8-yr old Yorkshire Terrier in Texas on 4/16 according to information from the National Animal Poison Control Center).
 - ii) There is no limitation on the dosage applied (such as spraying no more than so many seconds, or applying until the coat is only so moist). The exact wording of this should have been based on information from the registrant as to what constitutes an effective but safe dose and should have been correlated with actual dosage values (gms/kg) of product applied to dogs in studies.
 - iii) There is no limitation specified as to the frequency of treatments.
- 7. Toxicity studies on dogs were conducted. However, the maximum exposure was as a lX (use-direction) spray applied on a once-a-week basis over a period of 4 weeks. Adverse effects ("mild emesis" occurring 2 hours after exposure) were noted for two (out of 40) dogs in one of these studies. It is noted that a considerable number of dogs were used in more than one of these studies, that the only information provided regarding the individual dogs was breed, weight and hair coat (nothing as to age; 2 individuals in one study are, however, reported as "nursing puppies").

The dog toxicity studies (as well as the cat efficacy studies) were conducted by a laboratory for which the address is given (in one document)

None of these studies has a quality assurance statement, and there is no indication that this facility follows GLP regulations.

8. The Toxicology Branch has no record that any subchronic or chronic toxicity studies on DEET have been conducted on dogs. There is some concern regarding possible kidney and liver effects.

- 9. It is noteworthy that diarrhea is reported as a symptom in some incidents involving dogs. While no deaths were reported in the acute dermal rabbit LD50 study conducted (Leberco Testing, Inc.) on the concentrate, diarrhea (associated with weight loss) was reported from several animals, so that toxicity due to dermal absorption of the active cannot be ruled out. The only individual whose name appears on these reports is
- 10. A search through the available literature on DEET has failed to turn up any antidotal information, or anything other than "supportive treatment" for poisoning by this compound. This has made it difficult for the Toxicology Branch to respond to inquiries from veterinarians as to the appropriate procedure to follow.
- 11. The Toxicology Branch expresses reservations that additional toxicity testing as outlined in the draft review from IRB/TSS will expeditiously resolve the problems associated with these products.

- repels fleas
- repels ticks
- keeps your cat free of fleas and ticks for up to 21 days

N.N.Diethyl-m-tolusmide 8.55% Other Isomers Other Isomers
Cyano (3-phenoxyphenyl)
methyl-4-chloro-alpha(1-methylethyl) CTIVE INGREDIENTS:

EPA Reg. No. 2596-114 benzeneacetate

CAUTION: See back panel for additional Keep out of reach of children. precautionary statements. EPA Est. 2596-NJ-1

HARTZ BLOCKADE CAT FLEA AND TICK REPELLENT

repets fleas . repets ticks @ kills fleas . kills ticks @ repets mosquitoes keeps your pet flea and tick free for up to 21 days @ repels insects which

reduces incidence of itching and scratching

spray repelled fleas and ticks for up to 21 days. The spray may also be used on pets infested with fleas or ticks in order to kill these parasites and prevent Hartz Blockade Flea and Tick Repellent prevents these pests from actually inhabiting your cat. In laboratory and field tests, animals treated with this additional ones from inhabiting the pet.

DIRECTIONS FOR USE

It is a violation of Federal law to use this product in a manner inconsistent

quarters, and surrounding area should also be sprayed lightly until damp to help prevent reinfestation. Complete control can be obtained by affixing a Hartz 2 in 1® Reflecting Collar to the pet in addition to use of this spray. Under such circumstances, an occasional flea or tick may be seen on the animal. For more prolonged flea and tick control, the animal's bedding. to cover feet and tail. Do not spray in eyes. For best penetration of spray to the skin, direct spray against the natural lay of the fur to cause fluffing of the coat. Under normal circumstances, repeat spray every 21 days. In areas or hold container 6 to 10 inches from animal and spray over entire body. Be sure periods of heavy infestation it may be necessary to treat the cat more frequentto repel fleas and ticks, thereby reducing incidence of itching and scratching

Storage and Disposal Replace cap, wrap empty container and put in trash.

PRECAUTIONARY STATEMENTS

Hazards to Humans & Domestic Animals

CAUTION: Avoid inhalation of spray. Avoid spraying in eyes. It is good practice to wash hands after use. Bo not contaminate water or food.

Contents under pressure. Do not use or store near neat or open flame. Do not puncture or incinerate container. Exposure to temperatures above 130°F may cause bursting.

STATEMENT OF PRACTICAL TREATMENT
If in eyes. Rinse eyes with plenty of water. Call a physician if irritation persists.

Made and printed in U.S.A.

NOT REVIEWED

has been used that the pater β - β - β - β In Accordance with PR Noting 82-8:

ENCE OF POISONINGS IN SMALL ANIMALS

.mary problem of gastroinmes hemolysis. Paradichloess frequent but may be ataxia, and seizures.

icides. Amitraz (Mitaban. nded for dogs less than 3 es occasionally serious, or pisonings. Signs of serious occur in younger or very lude weakness, salivation, coma, and death. It is not ire to cause mild weakness arger dogs, and most such recover.

an Cyanamid), which cona heavily used fire ant action is poorly underdro is low, and toxicoses requent than exposures. nt is present in Combat ierican Cyanamid).

apparent effort to avoid ulations, crude citrus oil ted as "pet dips" that small print, followed by) large print) without accticidal. Such products " , and ataxia, depression, ilted even when label lowed. Some such forrawn from the market. not, however, be conng D-limonene (such as emicals, Inc.). Our exdips in cats suggest that commended concentraexposure, toxic effects on, hypothermia, and ts last minutes to seviage. In addition, scromassive overexposure sient signs sometimes d concentration.

l in origin, rotenone is fish but also to mamnly affected than dogs. re a result of topical ing dips or powders. in both dogs and cats osis or suspected toxariable but often inors, and dyspnea.

igone). Each year the regarding pennyroyal used as an insecticide eived in 1984 regardcosis consistently in-

described panting, chial sounds. These are compatible with reports of pennyroyal oil toxicosis in humans, the effects of which may also include disseminated intravascular coagulation, hepatic necrosis, abortion, and death.

REPELLENTS. Diethyltoluamide (DEET), found in many insect repellent products (such as Off, Deep Woods Off, Cutter Insect Repellent, and others [S. C. Johnson and Son and Cutter]), was associated with seven suspected cases of toxicosis or toxicosis calls in cats and four such calls in dogs. Naturally these occur during warm months and are a result of owner treatment of their pets. In dogs and, especially, cats, primary signs included seizures, tremors, and sometimes emesis. Ataxia or lethargy occasionally occurred...

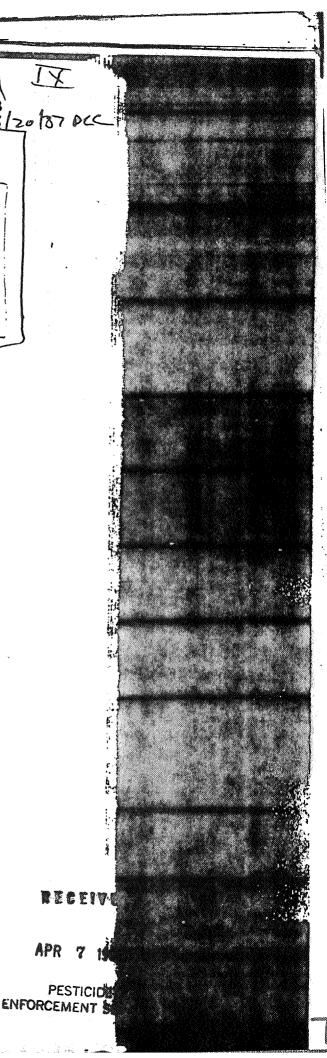
RODENTICIDES. The rodenticide class is the second most prevalent category of agents associated with calls to the NAPCC for two reasons: (1) Rodenticides are often placed in areas in which both rodents and pet animals are present, and (2) the NAPCC has an agreement with the manufacturers of brodifacoum so that the NAPCC telephone number is on packages containing this common anticoagulant (Talon and Havoc [ICI Americas and others]). Naturally, this causes us to receive a disproportionate number of calls concerning brodi-

Anticoagulant Rodenticides. The vast majority of anticoagulant rodenticide calls, regardless of the agent involved, were for exposure only. In the case of brodifacoum, the number of clinically serious poisonings resulting from exposure was undoubtedly minimized by virtue of access to appropriate antidotal information. It appears that dogs are more sensitive than cats to brodifacoum.

Owing to the extremely wide availability of anticoagulants, a significant number of toxicoses result from their use. Baits that contain bromadiolone, chlorophacinone, pindone, diphacinone, and others are occasionally consumed by small animals. Of these, it appears that diphacinone may cause the greatest number of poisonings (see page 159).

Strychnine. The convulsant strychnine continues to cause toxicosis, but the prevalence of this poisoning may be declining. Nevertheless, strychninecontaining baits are still available over the counter. During 1984, 26 calls pertaining to accidental and malicious poisonings of dogs were received. Only two such calls pertained to cats.

Fluoroacetate (Compound 1080). Fluoroacetate toxicosis is comparatively infrequent at the present time. This is a result of restrictions placed on the use of fluoroacetate-containing baits. Currently, fluoroacetate may be used as a rodenticide only by licensed exterminators. Depending on the outcome of legal battles continuing at the time of this writing, fluoroacetate may eventually be encountered as a result of its use in covote control in the form of "single lethal dose baits" and less often in toxic



CORE Grade/	DOC. NO.	Minimum 003891	Minimum 000490 	Minimum 000490 	Minimum 000490 	Minimum 000490	
T0X	Lategory	Not a sensitizer	II	_	2	——————————————————————————————————————	
	LD50, LC50, PIS, NOEL, LEL	No evidence for development of sensitization following a series of intradermal injections of 5% test material in corn oil. 0.05% DNCB (positive control) elicited sensitization reaction in all guinea pigs to which it was administered.	Minor corneal opacity with clearing lday 7 in 3/6 animals. Minor con- ljunctivitis with clearing by day 7 in 4/6 animals. (Unwashed eyes)	Minor corneal opacity present in 4/6 animals with clearing by day 21 (unwashed eyes). PIS = 30/110	No corneal opacity. No conjunctivi- tis.	No corneal opacity. Minor conjun- tivitis in 5/6 with clearing by day 3 in 2/5 animals.	
EPA Accession	No.	254183	244779	244779	244779	244779	
	Material	DEET35% Dimethyl phtha- late13% Di-n-propyl isocinchomero- nate1% N-Octyl bicyclo heptene dicar- lboximide1%	N,N-diethyl-m- toluamide . 15% (less viscous inerts)	N.N-diethyl-m- toluamide . 15% (more viscous inerts)	 Inert ingre- dients only 	Deet 15% (in corn oil) 	
Tox Chem No. 346 - U	Study/Lab/Study #/Date	ation - ies; 18/83	Primary eye irritation - rabbit; Biosearch, Inc.; #80-2264A; 12/31/80	Primary eye irritation - rabbit; Biosearch, Inc.; #80-2264A;	Primary eye irritation - rabbit; Biosearch, Inc.;	Primary eye irritation - rabbit; Biosearch, Inc.; #80-2137A; 10/20/80	

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	CORE Grade/ Doc. No.	Minimum 000825			Minimum 000825			Minimum 000825 			Minimum 000825			-
	TOX Category	11			II			⊢	·		I .	- <u></u>	حجب مست ہے۔۔۔	.
	Results: LD50, LC50, PIS, NOEL, LEL	PIS = 4.7 at 72 hrs (washed eye) PIS = 0.0 at 7 days (washed eye)	Corneal opacity present at day 3.	PIS = 0.8 at 72 hrs (unwashed) PIS = 0.0 at 7 days (unwashed)	PIS = 8.0 at 72 hrs (unwashed eyes) PIS = 0.0 at 7 days (unwashed eyes)	Corneal opacity present at day 3, bu cleared by day 7 for both washed and unwashed.	PIS = 4.6 at 72 hrs (washed eye) PIS = 0.0 at 7 day (washed eye)	pis = 13.8 at 72 hrs (unwashed eye) pis = 1.1 at 14 days (unwashed eye)	Corneal opacity present at day 14 for both washed and unwashed eyes.	PIS = 4.2 at 72 hrs (washed eye) PIS = 0.5 at 14 days (washed eye)	pis = 10.6 at 72 hrs (unwashed eye) pis = 0.0 at 7 days (unwashed eye)	Corneal opacity present at 72 hrs but cleared by day 7.		Page 2 of 4
EPA	Accession No.				- سيسا حسن حسد	والمحمد معتند وسين سنس				. حنید منب مب		 	· -	
Material		TECH 100%			75% solution DEET in dena-	tured EtOH	- · ·	 50% solution DEET in dena-	tured Ethanol 	حت جب حث		503 b) Insect Repellent 71.25% - DEET	25.0% - dena- tured Ethanol 3.25% - Other diethyl tola-	mides
117	dy/Lab/Study #/Date	Primary eye irritation - rabbit;	Dept. of Army; #75-51-0034-81; 1/1981		Primary eye irritation - rabbit;	Dept. of Army; #75-51-0031-81; 1/1981		Primary eye irritation - rabbit;	<pre>Dept. of Army; #75-51-0034-81; 1/1981</pre>		Primary eye irritation - rabbit:	Dept. of Army; #75-51-0034-81; 1/1981		

•	<u>-</u>			
	CORE Grade/ Doc. No.	Minimum 003891	Supplementary 003891	Minimum 003891 -
	TOX Category		(111)	> 1
	Results: LD ₅₀ , LC ₅₀ , PIS, NOEL, LEL	LD50(male) = 4550(3434-6029) mg/kg LD50(female)=5450(3759-7903) mg/kg Symptoms: hypoactivity, ptosis, ataxia, salivation, prostration, loss of righting reflex, bradypnea.	Three accidental deaths occurred at 2 g/kg (only dose tested).	10% mortality at 5.3 mg/L (actual concentration; only level tested); Nominal concentration = 39.1 mg/L; (4-hr exposure). Mass median diameter = 3.1 um, with 94% of particles < 9 um in diameter. Symptoms: ataxia, lethargy, salivaltion, respiratory distress. Inhalation LC50 (both sexes) > 5.3 mg/L (4-hr exposure; actual concentration).
ΡΡΔ	Accession No.	254183	254183	254183
-	Material	DEET22.95% Dimethyl phtha- late15.30% Di-n-propyl isocinchomero- nate1.275% N-Octyl bicyclo heptene dicar- boximide.1.275%	DEET22.95% Dimethyl phtha- late15.30% Di-n-propyl isocinchomero- nate1.275% N-Octyl bicyclo heptene dicar- looximide.1.275%	DEET22.95% Dimethyl phtha- late15.30% Di-n-propyl isocinchomero- nate1.275% N-Octyl bicyclo heptene dicar- boximide.1.275%
M NO. 340 - UEEI	ddy/Lab/Study #/Date	Acute oral LD50 - rat; Miles Laboratories; Report #28; 12/20/83	Acute dermal LD50 - rabbit; Miles Laboratories; Report #31; 12/20/83	Acute inhalation LC ₅₀ -rat; Bio-Research Lab. Ltd; Project No. 81774; 12/27/83

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CORE Grade/ Doc. No.	Minimum 003891	Minimum 003891	INVALID 001743	INVALID 001743
TOX Category	ΛΙ	111		
Results: LD50, LC50, PIS, NOEL, LEL	PDIS =0.21 (24-hr occluded exposure)	All eyes (wahsed and unwashed) had some irritation. Some eyes had corneal involvement. All eyes were clear by day 7.	Serious inconsistencies in the reported data.	Serious inconsistencies in the reported data.
EPA Accession No.	254183	254183	252568	252568
Materia]	DEET22.95% Dimethyl phtha- late15.30% Di-n-propyl isocinchomero- nate1.275% N-Octyl bicyclol heptene dicar- boximide.1.275%	DEET22.95% Dimethyl phtha- late15.30% Di-n-propyl isocinchomero- nate1.275% N-Octyl bicyclo heptene dicar- boximide.1.275%	Deet 100% 252568	Deet 20% Ethanol 80%
	Primary dermal irrita- tion - rabbit; Miles Laboratories; Report #27; 12/20/83	Primary eye irritation- rabbit; Miles Laboratories; Report #18; 11/18/83	Primary eye irritation - rabbit; Medical College of Wisconsin; 1983	Primary eye irritation - rabbit; Medical College of Wisconsin; 1983

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