

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

OFFICE OF PREVENTION, PESTICIDES AND TOXIC SUBSTANCES

Scott Hathorn III Regulatory & EHS Manager KMG-Bernuth, Inc. 9555 W Sam Houston Pkwy S. Suite 600 Houston, TX 77099

MAR 1 1 2009

SUBJECT:

Applications for Pesticide Notification – Labeling Changes per PR Notices 98-10

Rabon® 97.3 Oral Larvicide

EPA Reg. No. 61483-47

Add Instructions in the Event of Separation of the Supplemental Mixing and Use Instructions from

Product Container

PermectrinTM Fly & Louse Dust

EPA Reg. No. 61483-67

Add Pest Graphics

Applications Dated October 28, 2008

Dear Mr. Hathorn III:

The Agency is in receipt of your Applications for Pesticide Notification under Pesticide Registration Notices (PRNs) 98-10 for the above products. The Registration Division (RD) has conducted a review of these requests for their applicability under PRNs 98-10 and finds that the actions requested fall within the scope of these PRNs. The labels submitted with the applications have been date-stamped "Notification" and will be placed in our records.

If you have any questions, please call me directly at 703-305-6249 or Terri Stowe of my staff at 703-305-6117.

Sincerely,

Linda Arrington

Notifications & Minor Formulations Team Leader

Registration Division (7505P)

Office of Pesticide Programs

Please read instructions on reverse before	re completing form		Form /	Approved OMI	3 No. 2070-0080. Aj	proval	expires 2-28-98		
Unite	d States			Regist	ration	OP	P Identifier Number		
EPA Environme	ncy		Amend	ment					
Washington	_	X	Other						
	Application for Pe	sticide - Section 1	1						
1. Company/Product Num	ber	2. EPA Product N	/lanaç	jer		3.	Proposed Classification		
61483-47		George LaRocca			_ _				
Company/Product (Nar Rabon® 97.3 Oral Larvi	PM#	13			X None Restricted				
5. Name and Address of		de 6. Expedited Review		ordance wil	th FIFRA Section	n 3(c)	(3)(b)(i),		
KMG-Bernuth	• • • • •	my product is similar of							
	Houston Pkwy S. Ste 60	0 EPA Reg. No.							
Houston, Tex		Draduat Nama							
X Check if t	this is a new address	Product Name							
	Section -	- II							
Amendme	nt - Explain Below.	· · · · · · · · · · · · · · · · · · ·		Final prin	ted labels in res	ponse	to		
				. ~ ^	etter dated July	27, 20	005		
Resubmis	ır		"Me Too"	le Too" Application					
dated X Notification			Other - F	xplain below.					
The purpose of this notific		pecific identification i	nform			enta	I Mixing and Use		
Instructions. KMG-Bernu									
become separated from t	he product container an	d if so, would not ha	ive su	ifficient i	dentification	n to :	reference the end user		
back to the container. S	pecifically added is the	active ingredient stat	emen	t, the EP	A Reg. No.	, the	EPA Est. No.,		
the Keep Out of Reach o	of Children statement an	d the Signal Word.							
-		•							
	Section	- 111							
 Material This Product V 									
Child-Resistant Packaging	Unit Packaging		Water Soluble Packaging			2. T	2. Type of container		
Yes No	Yes		Yes		Metal				
* Certification must	 	. per	if "Yes"		No. per	+	Plastic Glass		
he submitted		ainer	Package Wt		container	Paper			
							Other (specify		
3. Location of Net Contents Infor		4. Size(s) Retail Conta	ainer		5. Location of	of Lab	el Directions		
Label 6. Manner in Which Label is Affix	Container			Lithograph	L	T	Stenciled		
o. Mainter in Willon Label is Allix		Paper glued			Other				
	Section	- IV							
1. Contact Point (Complete items	directly below for identification	of individual to be contact	ed, if n	ecessare, to	process this a	pplica	tion.)		
Name		Title	•			Tele	phone No. (Include Area Code)		
Mr. Scott Hathorn III Regulatory & EHS Manager					(713) 600-3819				
Logitify that the statements I h	Certification have made on this form and all a	ettachmente therete ere true	20011	ata		6. 1	Oate Application Received		
	that any knowingly false or misic						(Stamped)		
by fine or imprisonment or both		July 50 pt					00000		
	· · · · · · · · · · · · · · · · · · ·						0 000000		
2. Signature		3. Title							
De Holulat	P D'UN	Pogulator: 9F116	Regulatory &EHS Manager			0000			
NI MILIMA	11. 0 -1	regulatory &EHS	o Mar	ager	Ļ	د د د	. (.		
4. Typed Name		5. Date				٠,٠	,		
		October 28, 2008	8		(,	(() ((, (0 66 6		



October 28, 2008

Document Processing Desk (NOTIF)
Office of Pesticides Programs (7504P)
U. S. Environmental Protection Agency
Ariel Rios Building
1200 Pennsylvania Avenue, N. W.
Washington, D.C. 20460

Re:

RABON® 97.3 Oral Larvicide

EPA Reg. No. 61483-47

Dear Mr. Larocca:

This notification is a resubmission of the final printed Supplemental Mixing and Use Instructions for the subject product. This resubmission is being made to alleviate the possibility of misidentification of the instructions in the event of separation from the product container. Please see Section II of the included application form for the specifics added to the instructions.

Specifically enclosed in support of this notification are the following documents:

- 1. EPA Application Form 8570-1 dated October 28, 2008
- 2. Three (3) copies of the revised Supplemental Mixing and Use Instructions.

We are requesting an expedited review of this submission as well as a stamped copy of the approved label for use in submitting registration requests to the various states.

"This notification is consistent with the provisions of PR Notice 98-10 and EPA regulations at 40 CFR 152.46, and no other changes have been made to the labeling or the confidential statement of formula of this product. I understand that it is a violation of 18 U.S.C. Sec. 1001 to willfully make any false statement to EPA. I further understand that if this notification is not consistent with the terms of PR Notice 98-10 and 40 CFR 152.46, this product may be in violation of FIFRA and I may be subject to enforcement action and penalties under sections 12 and 14 of FIFRA."

Thank you for your assistance and if you have any questions, please contact me.

fe7 - Scott Hathorn III

Regulatory & EHS Manager

On Youth R. O'Ned

Ph. 713-600-3819

ONLY FOR USE IN CATTLE FEEDS

NOTIFICATION

MAR 1 1 2009

Supplemental Mixing and Use Instructions

EPA Est. No. 61483-KS-01

Active Ingredient:	By Weight
Tetrachlorvinphos (CAS No. 22248-79-9)	97.3%*
Other Ingredients:	2.7%*
Total:	100.0%*
*RABON Technical – 442 grams per pound	

Keep Out of Reach of Children

CAUTION

EPA Reg. No. 61483-47



Manufactured by: KMG-Bernuth, Inc. 9555 W. Sam Houston Pkwy. S. Suite 600 Houston, Texas 77099 800-322-8177

000000

000000 800800

00000 42:326122270/00-0809

EPA 091807 Corr FPL

RABON 97.3

Oral Larvicide ONLY for Use in Cattle Feeds

Rabon Oral Larvicide prevents the development of Horn Flies, Face Flies, House Flies and Stable Flies in the manure of treated cattle. The following information is supplemental to the product label. Prior to use of this product, also read and observe the precautions on the label affixed to the product container.

DESCRIPTION

RABON is the registered trade name for Tetrachlorvinphos: 2-chloro-1 (2, 4, 5-trichlorophenyl) vinyl dimethyl phosphate. RABON Oral Larvicide is a specially prepared granulated material designed to provide optimum larvicidal activity in the manure. RABON 97.3 Oral Larvicide is available to the feed manufacturer for formulating larvicidal feeds.

ACTION

When fed to cattle, RABON Oral Larvicide passes through the digestive system into the animal's manure where it kills fly larvae on contact shortly after fly eggs hatch. By preventing larval development, RABON Oral Larvicide helps to control adult fly populations.

Manure from treated cattle will remain larvicidal up to six weeks; manure older than six weeks will not support fly development unless it becomes wet or contaminated with fresh manure.

TOXICOLOGY

The toxicology of RABON Oral Larvicide has been investigated in extensive field and laboratory studies in both domestic animals and wildlife. Cattle of both sexes, various ages and breeds, and maintained under a variety of management conditions, have been treated with many times the recommended dose. Milk production, reproduction, growth, and feed efficiency were not adversely affected. RABON Oral Larvicide can be used in conjunction with other organophosphate insecticides or pyrethroid compounds.

Test animals refused to eat excessive amounts of RABON Oral

Larvicide. Care must be taken when feeding RABON Oral Larvicide to newborn calves. Adverse reactions in calves are not seen when RABON Oral Larvicide is fed in the grain portion of the diet, but intoxication can occur if excessive amounts are fed through the milk.

Overdosing with this compound will result in only slight inhibition of the enzyme cholinesterase within the nervous system and the neuromuscular junction. Intoxication can be reversed by the administration of atropine.

PALATABILITY

RABON Oral Larvicide is palatable to cattle when used as directed.

EFFECT ON THE ENVIRONMENT RABON Oral Larvicide does not affect beneficial insects such as dung beetles or other insect predators that normally inhabit the manure. The manure from treated cattle may be used immediately as fertilizer.

To prevent pollution of the environment, good sanitation practices should always be followed when disposing of animal wastes.

AREA-WIDE CONTROL

RABON Oral Larvicide prevents the development of horn flies, face flies, house flies, and stable flies in the manure of treated cattle but does not affect existing adult flies. Since flies tend to migrate from farm to farm, the use of a feed additive larvicide should be considered as only a part of the total fly control program. Periodic spraying of buildings or animals with other insecticides may be necessary in order to control invading adult flies.

FEEDING AND MANAGEMENT

Start feeding RABON Oral Larvicide early in the spring before flies begin to appear and continue feeding throughout the summer and into fall until cold weather restricts fly activity. The proper feeding period will vary with climate and should be determined by the emergence date of flies in previous years for your area.

Rations containing RABON Oral Larvicide may be fed up to slaughter and to lactating dairy cows without withholding the milk from market during or after treatment.

RABON Oral Larvicide should be used in conjunction with other good management and sanitation practices. All potential fly breeding material such as manure, old hay, and silage which contains overwintering fly pupae should be removed from the premises. Manure should not be allowed to accumulate around barns, fences, or under feed bunks during the fly breeding season.

When starting a feeding program during the fly season, it is desirable to use other control measures to reduce the population of existing adult flies.

In some cases, supplemental fly control measures may be needed in and around dry lots, calf pens, and barns to control adult house flies and stable flies which can breed not only in cattle manure but in other decaying vegetable matter or silage on the premises.

RECOMMENDED FEEDING LEVEL

For effective fly control, it is important to insure that all cattle on the premises receive adequate levels of RABON Oral Larvicide on a daily basis. The recommended feeding level of RABON Oral Larvicide to cattle is 70 mg per 100 pounds of body weight daily. The amount of RABON Oral Larvicide consumed by individual animals on a daily basis may vary, but fly larvae control will not be affected. A practical feeding regimen can be planned whereby a single feed cap before to all animals within a like group. Examples are illustrated below.

Growing-Finishing Beef Cattle

A single complete feed containing 26.4 mg of RABON per pound of feed can be fed to all steers and/or heifers, weighing from 400 to 1400 pounds, within a group. As long as the daily feed consumption of the larvicidal feed approximates that shown in the chart (See Figure 1), the cattle will be receiving the larvicide at an acceptable level.

Lactating Dairy Cows

A single concentrate feed containing 66 mg RABON per pound of feed fed along with roughage can be fed to all lactat-

ing cows within the herd. As long as the daily feed consumption of the larvicidal feed approximates that shown in the chart (See Figure 2), the cows will be receiving RABON at an acceptable

Hand-Fed Beef or Dairy Cattle

A common practice in many cattle operations is to hand-feed a supplement at a given level per head daily. A single supplement which contains 792 mg RABON per pound of supplement and which is fed at the rate of 1 pound per head daily can be fed to cattle

weighing between 400 to 1200 pounds. Cattle weighing between 1200-1700 pounds should be fed this supplement at the rate of 1 ½ pounds per head daily.

DIRECTIONS FOR MIXING LARVICIDAL FEEDS

RABON Oral Larvicide will mix uniformly in cattle feeds when good blending procedures are followed. Blending studies performed in typical feed mills indicated that mixer coefficients of variation of 2 to 8 percent can be achieved. Directions for mixing various cattle feeds with RABON Oral Larvicide are given in Table 1.

Feeds prepared using this product should not be pelleted unless tests are conducted to assure adequate RABON levels after pelleting. Do not mix this product with feeds containing predominantly pellets due to particle size differences and potential segregation. This product can be used in liquid feed supplements (LFS) provided recommended guidelines are followed.

STABILITY

Under normal warehouse conditions, RABON 97.3 Oral Larvicide is stable for a minimum of 2 years.

RABON Oral Larvicide when mixed with several typical cattle rations in stability tests, was found to be stable (<10% decomposition) for up to 3 months in complete feeds, for up to 6 months in protein supplements, and for up to 1 vear in mineral mixtures when stored under normal conditions. Storage of feedstuffs containing RABON Oral Larvicide at elevated temperatures (ca 100 °F) for an extended period does have a detrimental effect on the stability of RABON; therefore, complete feeds, protein supplements, and liquid feeds stored under such conditions should be fed within 4-8 weeks of manufacture.

ANALYTICAL METHODS

Sampling procedures, analytical procedures, and standards used for determining RABON Oral Larvicide in various feed products are available upon request from KMG-Bernuth, Inc.

Figure 1.
Daily Consumption of Complete
Ration (Containing 26.4 mg
Rabon per pound
of feed) Required
for Fly Larvae
Control

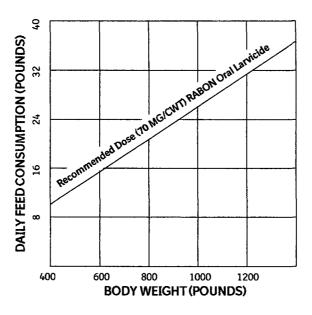


Figure 2.
Daily Consumption of Dairy Feed (Containing 66 mg Rabon per pound of feed)
Required for Fly Larvae Control

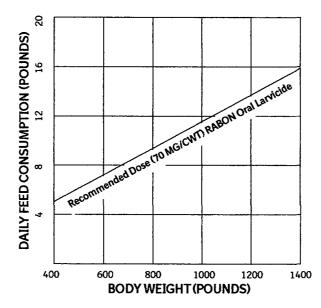


TABLE 1. Mixing Directions in Dry Feed								
			ON in ement	Ratio Supplement		evel of in Feed	Pounds RABON 97.3% Oral	
Feed Product	Daily Feeding Rate	MG/LB	%	to Feedstuffs	MG/LB	%	Larvicide/ Ton Product	
Complete Feed			N/A	N/A	26.4	0.0059	0.12	
	N/A	132	0.029	1:4	26.4	0.0059	0.6	
Supplement or Premix for	N/A N/A	264 528	0.059 0.12	1:9 1:19	26.4 0.0059 26.4 0.0059		1.2 2.4	
Preparing Complete Feeds	N/A	1056	0.24	1:39	26.4	0.0059	4.8	
	N/A	2112	0.47	1:79	26.4	0.0059	9.6	
Concentrate Fed with Roughage	1.0 lbs./cwt.	N/A	N/A	N/A	66	0.015	0.3	
	N/A	220	0.070	4.4	66	0.045	4.5	
Supplement or	N/A N/A	330 660	0.072 0.15	1:4 1:9	66 66	0.015 0.015	1.5 3.0	
Premix for Preparing	N/A	1320	0.13	1:19	66	0.015	6.0	
Concentrate	N/A	2640	0.59	1:39	66	0.015	12.0	
Feeds	N/A	5280	1.17	1:79	66 0.015		24.0	
	2.0 lbs./head	396	0.087	N/A	N/A	N/A	1.8	
Supplement for	1.5 lbs./head	528	0.12	N/A	N/A	N/A	2.4	
Hand Feeding	1.0 lbs./head	792	0.18	N/A	N/A N/A		3.6	
	0.5 lbs./head	1584	0.35	N/A	N/A	N/A	7.2	
	4.0 oz./head	3168	0.71	N/A	N/A	N/A	14.4	
	3.0 oz./head	4224	0.93	N/A	N/A	N/A	19.2	
8.4' 1.8.4'	2.0 oz./head	6336	1.40	N/A	N/A	N/A	28.8	
Mineral Mix	1.0 oz./250 lbs.	2700	0.60	· N/A	N/A N/A N/A		12.3	
	1.0 oz./500 lbs.	5400	1.19	N/A	N/A	N/A	24.6	
	1.0 oz./750 lbs.	8100	1.79	N/A	N/A	N/A	36.9	

Note: Calculations presented in this table serve as guidelines in preparing feeds which contain RABON Oral Larvicide. Some calculation inconsistencies occur in the table due to rounding. A head weight of 1165 lbs. was used for these calculations. N/A = Not Applicable

CHARACTERISTICS OF LIQUID FEED SUPPLEMENTS (LFS)

A great diversity of suspension formulations exist in the liquid feed industry. The degree of variability in formulation content is dependent upon which market the finished feed product is going to be used (i.e., pasture beef, dairy cattle or the feedlot market). Variation in moisture, sugar, mineral and protein content as well as the intended consumption rate, all affect the composition of the finished product. Therefore, no two LFS possess the same physical and chemical characteristics. One of the most important physical characteristics with respect to ROL in LFS is viscosity, which can vary dramatically. Viscosity of a given medium plays an important role in keeping insoluble particulate matter such as ROL in suspension.

In order to successfully use ROL in LFS, two criteria must be met. First, the ROI

CRITERIA FOR ROL USE IN LFS

two criteria must be met. First, the ROL must remain positionally stable in the LFS with no settling during the feedout period (6-8 weeks). Second, the ROL must be chemically stable in the LFS.

SUSPENDABILITY OF ROL IN LFS

Kelflo™ is a high quality xanthan gum polysaccharide product that has been specially formulated as a suspending and stabilizing agent in LFS. Solutions of Kelflo xanthan gum exhibit pseudoplastic (or shear-thinning) flow behavior. That is, stored LFS have excellent suspension stability. When shear is applied (mixing, pumping, spraying, transport, lick wheel movement), the viscosity decreases, resulting in a freeflowing liquid. When shear is removed, the original at-rest viscosity is restored so that suspension stability is maintained. Xanthan gum is routinely used at 3 lb/ton LFS by some manufacturers to suspend drug products.

Laboratory studies have shown that hydrated xanthan gum at various concentration levels will suspend ROL in LFS. As a general rule, LFS with viscosities of 400–600 centipoise (cP, Brookfield viscometer at 20 rpm, spindle No. 3 for 2.5 minutes at 68–72°F) require 3 lb xanthan gum for LFS to provide positional stability of ROL.

This amount of xanthan gum increases the viscosity of the final product above 1900 cP. Xanthan gum must be fully hydrated prior to the addition of ROL. In the laboratory, xanthan gum requires 24 hours to become hydrated. For this reason, we recommend that xanthan gum be added to the LFS at manufacture, not at the distributor or consumer level.

For LFS with low viscosities (i.e., 30-55 cP), 9-12 lb xanthan gum/ton LFS would likely be required to suspend ROL. However, the cost/ton of LFS would most likely be prohibitive.

GUIDELINES

The following steps are recommended for the successful use of ROL in LFS:

- 1. The LFS manufacturer should determine the viscosity of the liquid product being considered for ROL inclusion. The viscosity should be performed using a Brookfield viscometer at 20 rpm, spindle No. 3 for 2.5 minutes at 68-72°F. The viscosity reading will establish the feasibility of adding ROL to the LFS.
- For LFS with viscosity readings of 400-600 cP at 20 rpm, the addition of 3 lb xanthan gum/ton LFS appears to adequately suspend ROL.
- For LFS with very low viscosities, 9-12 lb xanthan gum/ton LFS will likely suspend ROL. Manufacturers interested in suspending ROL in such low viscosity LFS should contact KMG-Bernuth, Inc. for technical assistance or conduct suspendability studies themselves.
- Addition of xanthan gum to the LFS must be conducted during manufacture in order to ensure full hydration of the gum prior to the addition of ROL.
- ROL and xanthan gum should not be added to the LFS at the same time. The ROL will settle out.
- ROL and xanthan gum should not be added to the LFS at manufacture. However, ROL could be added

to LFS (previously charged and hydrated with the proper level of xanthan gum) at the distributor level provided adequate mixing of the ROL is accomplished.

- 7. ROL remains stable chemically when mixed and stored in products with low pH values (6 or less).
- Field trials to determine the positional and chemical stability of ROL in LFS should be conducted. Dating of the product (normally 6-8 weeks) will be based on the results of the field trial.
- 9. LFS containing ROL should be fed out in 6-8 weeks.
- 10. Lick tanks should be emptied prior to refilling to ensure the maximum dating of the product. Chemical stability and efficacy of the product in the field will be severely compromised if fresh product is mixed with outdated material.
- 11. LFS with ROL must be accompanied with end-use labeling directions.
- 12. ROL can be added to LFS containing drug products.

DIRECTIONS FOR MIXING IN LIQUID FEED SUPPLEMENTS

This product is a specially prepared granulated material designed for use in cattle Liquid Feed Supplements (LFS). The proper positional and chemical stability of ROL in liquid feeds is dependent upon acceptable physical characteristics and proper blending technique. Depending on the viscosity of the liquid feed, it may be necessary to add xanthan gum.

The Liquid Feed Supplement (LFS) plus xanthan gum must be fully hydrated prior to the addition of the RABON Oral Larvicide.

RABON Oral Larvicide should be fed in Liquid Feed Supplements (LFS) for cattle (including lactating dairy cattle) to supply the recommended feeding level of 70 mg of this product per 100 pounds of body weight daily.

of LFS to be fed per animal per day. Use Surement, for determining the proper Table 2 below as a guide, depending on mixing rate. the most convenient method for mea-

Pounds LFS	Cattle Weight	600 lb.	800 lb.	1000 lb.	1200 lb.	1350 lk
Consumed/Day 1.0		850 g.	1134 g.	1418 g.	1701 g.	1928 (
1.5	Grams of Rabon 97.3 Oral Larvicide/TON of LFS ^a	624 g.	850 g.	1048 g.	1247 g.	1418
2.0		425 g.	567 g.	709 g.	850 g.	964
			- 1			2
Pounds LFS Consumed/Day	Cattle Weight	600 lb.	800 lb.	1000 lb.	1200 lb.	1350 II
1.0	Fluid Ounce Equivalents ^b of Rabon 97.3 Oral Larvicide/TON of LFS ^a	34 oz.	45 oz.	56 oz.	67 oz.	76 o
1.5		25 oz.	34 oz.	42 oz.	50 oz.	56 o
2.0		17 oz.	23 oz.	28 oz.	34 oz.	38 o