|  | AN ADDAMACHINAL DOOTCOYLON ACCAICY   |  | DATE OF ISSUANCE  |
|--|--|--|---|
|  | NVIRONMENTAL PROTECTION AGENCY<br>DFFICE OF PESTICIDES PROGRAMS<br>REGISTRATION DIVISION (75-767)<br>WASHINGTON, DC 20460  | E.4439 - 1<br>TERM OF ISSUANCE   | AUG <u>9 3 1994</u>   |
| NOTICE   | OF PESTICIDE   | NAME OF PESTICIDE PRO  | DUCT  |
| (Unde  | NEREGISTRATION<br>er the Federal Insecticide, Fungicide,<br>ad Rodenticide Act, as amended)  | Mole Med   |   |
| AME AND ADI  | DRESS OF REGISTRANT (Include ZIP code)   |  |   |
| <b>F</b>   |  | т 🔅  | 5/161227 141  |
| P.0  | e Med Inc.<br>. Box 333  |  | 5458168 146   |
| Auro<br>L  | ora, Indiana 47001   | L  |   |
| ubmitted to  | es in labeling formula differing in substance f<br>and accepted by the Registration Division prio<br>s refer to the above U.S. EPA registration num  | r to use of the label in comm  | ion with this registration must be<br>erce. In any correspondence on this   |
|  | of information furnished by the registrant, the nsecticide, Fungicide, and Rodenticide Act.  | above named pesticide is he  | reby Registered/Reregistered under  |
| A copy of the  | labeling accepted in connection with this Re-  | gistration Reregistration is   | returned herewith.  |
| cide in accor<br>Act is not to                             | e environment, the Administrator, on his motio<br>dance with the Act. The acceptance of any na<br>be construed as giving the registrant a right to   | me in connection with the re   | gistration of a product under this  |
| cide in acco<br>Act is not to<br>by others.                | dance with the Act. The acceptance of any na   | onally registered  | gistration of a product under this<br>or to its use if it has been covered<br>in accordance   |
| cide in acco<br>Act is not to<br>by others.                | dance with the Act. The acceptance of any na<br>be construed as giving the registrant a right to<br>This product is uncondition  | onally registered<br>ne year, provided   | gistration of a product under this<br>or to its use if it has been covered<br>in accordance<br>that you:  |
| cide in acco<br>Act is not to<br>by others.<br>wit]        | dance with the Act. The acceptance of any na<br>be construed as giving the registrant a right to<br>This product is uncondition<br>h FIFRA sec. 3(c)(5), for on<br>Make the following changes  | onally registered<br>ne year, provided<br>s to your label p  | gistration of a product under this<br>or to its use if it has been covered<br>in accordance<br>that you:<br>rior to releasing   |
| cide in acco<br>Act is not to<br>by others.<br>wit]        | dance with the Act. The acceptance of any na<br>be construed as giving the registrant a right to<br>This product is uncondition<br>h FIFRA sec. 3(c)(5), for on<br>Make the following changes<br>the product for shipment:   | onally registered<br>ne year, provided<br>s to your label p<br>" to "Mole Repelle  | gistration of a product under this<br>or to its use if it has been covered<br>that you:<br>rior to releasing<br>ent".   |
| cide in acco<br>Act is not to<br>by others.<br>wit]        | dance with the Act. The acceptance of any ma<br>be construed as giving the registrant a right to<br>This product is uncondition<br>h FIFRA sec. 3(c)(5), for on<br>Make the following changes<br>the product for shipment:<br>1. Change "Mole Repellant  | onally registered<br>ne year, provided<br>to your label p<br>to "Mole Repelle<br>statement as folle  | gistration of a product under this<br>or to its use if it has been covered<br>that you:<br>rior to releasing<br>ent".   |
| cide in acco<br>Act is not to<br>by others.<br>wit]        | dance with the Act. The acceptance of any ma<br>be construed as giving the registrant a right to<br>This product is uncondition<br>In FIFRA sec. 3(c)(5), for on<br>Make the following changes<br>the product for shipment:<br>1. Change "Mole Repellant<br>2. Revise the ingredient s<br>Active Ingredient  | onally registered<br>ne year, provided<br>to your label p<br>to "Mole Repelle<br>statement as folle  | gistration of a product under this<br>or to its use if it has been covered<br>in accordance<br>that you:<br>rior to releasing<br>ent".<br>ows:<br>. 66.0%   |
| cide in acco<br>Act is not to<br>by others.<br>wit]        | <pre>dance with the Act. The acceptance of any na<br/>be construed as giving the registrant a right to<br/>This product is uncondition<br/>h FIFRA sec. 3(c)(5), for on<br/>Make the following changes<br/>the product for shipment:<br/>1. Change "Mole Repellant<br/>2. Revise the ingredient s<br/>Active Ingredient<br/>Castor Oil (USP)<br/>Inert Ingredients<br/>3. Because Mole-Med is Tom<br/>exposure, no statement</pre>                           | onally registered<br>ne year, provided<br>s to your label p<br>" to "Mole Repell<br>statement as foll<br><br>xicity Category I<br>is needed for th<br>ecause Mole-Med i                        | <pre>gistration of a product under this<br/>or to its use if it has been covered<br/>that you:<br/>rior to releasing<br/>ent".<br/>ows:<br/>. 66.0%<br/>. 34.0%<br/>V for acute oral<br/>is route of<br/>s Toxicity</pre>                       |
| cide in accor<br>Act is not to<br>by others.<br>with<br>A. | <pre>dance with the Act. The acceptance of any ma<br/>be construed as giving the registrant a right to<br/>the FIFRA sec. 3(c)(5), for on<br/>Make the following changes<br/>the product for shipment:<br/>1. Change "Mole Repellant<br/>2. Revise the ingredient<br/>Castor Oil (USP)<br/>Inert Ingredients<br/>3. Because Mole-Med is To<br/>exposure, no statement<br/>exposure. However, be<br/>Category III for Prima:</pre>                            | onally registered<br>ne year, provided<br>s to your label p<br>" to "Mole Repell<br>statement as foll<br><br>xicity Category I<br>is needed for th<br>ecause Mole-Med i                        | <pre>gistration of a product under this<br/>or to its use if it has been covered<br/>that you:<br/>rior to releasing<br/>ent".<br/>ows:<br/>. 66.0%<br/>. 34.0%<br/>V for acute oral<br/>is route of<br/>s Toxicity<br/>ent is needed for</pre> |
| cide in accor<br>Act is not to<br>by others.<br>with<br>A. | <pre>dance with the Act. The acceptance of any na<br/>be construed as giving the registrant a right to<br/>This product is uncondition<br/>h FIFRA sec. 3(c)(5), for on<br/>Make the following changes<br/>the product for shipment:<br/>1. Change "Mole Repellant<br/>2. Revise the ingredient s<br/>Active Ingredient<br/>Castor Oil (USP)<br/>Inert Ingredients<br/>3. Because Mole-Med is To:<br/>exposure, no statement<br/>exposure. However, be</pre> | onally registered<br>ne year, provided<br>s to your label p<br>" to "Mole Repelle<br>statement as follo<br><br>xicity Category I<br>is needed for th<br>ecause Mole-Med i<br>ry Skin, a statem | <pre>gistration of a product under this<br/>or to its use if it has been covered<br/>that you:<br/>rior to releasing<br/>ent".<br/>ows:<br/>. 66.0%<br/>. 34.0%<br/>V for acute oral<br/>is route of<br/>s Toxicity<br/>ent is needed for</pre> |

c. Make the following change in the "DIRECTIONS FOR USE":

Change "After treatment ... an additional 25 minutes." to "After treatment, soak in with water using hose or sprinkler for about 25 minutes. Avoid using excess water that may flow off turf into streams, ponds, gutters or storm sewers."

7. For a product size intended for domestic, outdoor use change your Storage and Disposal statement as follows:

STORAGE AND DISPOSAL

- STORAGE: Store only in original container, in a dry place inaccessible to children and pets.
- DISPOSAL; Do not reuse empty container. Rinse thoroughly. Securely wrap in newspaper and discard in trash.
- 8. Add the following to your label:

EPA Registration No. 64439-1.

- B. Within one year of registration, or August 15, 1995, you need to submit data to answer the questions raised by the Dudderar and Elshof study (MRID 431216-04), as follows:
  - 1. The study was greatly curtailed due to limited funding. As no monitoring of mole activity in untreated areas took place, we concur with the authors that the

"study cannot link ceased activity directly to product application."

On April 28, 1994, Dr. Dudderar informed a member of my staff that he has students conducting additional trials with MOLE-MED this Spring and that these studies were expected to be completed within a month. As these trials would include the monitoring of untreated areas, they would seem to be designed better than the study run in 1993. We feel that at least some sham-treated (water only) areas should be monitored as flooding is occasionally mentioned as a method for controlling (or moving) voles. We also have suggested that small tests of the effects of MOLE-MED on earthworms and other prey be run to indicate whether the product's claimed effects on moles could be mediated indirectly.

As soon as we receive the results of these studies, we will inform you of the results. If those tests are



satisfactory, you need do no more testing. If those tests are not conducted or are otherwise unsatisfactory, we will provide you with additional guidance concerning testing. In this case you will be responsible for supplying additional data by the **August 15, 1995**, date. 39 4

2. The item "FAXed" to us on March 21, 1994, concerning the source of inspiration for this product was interesting. If this approach truly has merit for mole control, as Mrs. Bjerstedt reported, it should be possible to demonstrate the extent and duration of such effects in appropriately designed research studies.

If you do not satisfy this requirement for acceptable efficacy data by August 15, 1995, the Agency will automatically cancel this registration, effective on that date.

C. Submit one copy of your final printed labeling before you release the product for shipment. Refer to the A-79 Enclosure for a further description of final printed labeling.

This registration will be subject to cancellation in accordance with FIFRA sec. 6(e) if you do not comply with these conditions. Your release for shipment of the product constitutes acceptance of these conditions.

A stamped copy of the label is enclosed for your records.

RAF

Robert A. Forrest Product Manager (14) Insecticide-Rodenticide Branch Registration Division (H7505C)

Enclosures: 1) Stamped label 2) A-79 Enclosure

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LAWN PROTECTION



## COMPOSITION

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|------------|---------------|--------|-----|-------|-------|
| Oil of Pic | 66 <b>%</b> , |        |     |       |       |
| B->tegra   | dable Sulta   | clants |     | •••   | 143,  |
| nort       |               |        |     |       | . 20% |
| Keel       | o out of      | reach  | ofe | hildr | en    |

# WARNING

# AVISO

PRECAUCION AL USUARIO:Si usted no lee ingles, no use este producto hasta que le eliqueta haya sido explicado ampliamente.

STATEMENT OF PRACTICAL TREATMENT: IF IN EYES. Flush with plenty of water, get medical attention IF SWALLOWED: Drink promptly a large quantity of milk, eqg whites, gelatin solution, or, if those are not available, drink large quantities of water: Avoid alcohol

> She back panel for additional precautionary statements

**DELCONTENTS 32 OZ** 

MOLE-MED P.O. Box 333 Aurora, Indiana 47001 For repelling moles in thems

SHAKE WELL BEFORE USING PRECAUTIONARY STATEMENTS HAZARDS TO HUMANS AND DOMESTIC ANIMALS

WARNING: Cause substantial but temporary eye injury: -Do not get in eyes or on clothing. Wear salety glasses May be harmful il swallowed. Prolonged or fraquent repeated skin contact with this product may cause allergic. akin reactions in some individuals. Wash thoroughly with soap and water after handling.

#### **DIRECTIONS FOR USE**

It is a violation of Federal law to use this product in a manner inconsistent with its labeling.

USE RESTRICTIONS: For repeiling eastern moles and Townsend's moles from lawns.

MIXING DIRECTIONS: Mix with water at a rate of one ounce of MOLE-MED per gallon of water. Use the DILU-TION TABLE below to determine the amount of mixture to prepare for the area that you intend to treat. SHAKE MOLE-MED CONTAINER WELL BEFORE MIXING.

### DILUTION TABLE

| Amount of<br>MOLE MED | Amount of<br>Water | Area to be<br>Covered |
|-----------------------|--------------------|-----------------------|
| 1 oz.                 | 1 gal.             | 312 sq. ft.           |
| 2 oz.                 | 2 gal.             | 624 sq. ft.           |
| 16 oz.                | 16 gal.            | 5,000 sq. ft.         |
| 32 oz.                | 32 gal.            | 10,000 sq. ft.        |

LOCATING MOLES: The presence of moles may be Indicated by a network of surface ridges in the turf or by a series of conical mounds of earth pushed up from deep burrows. Treated all areas which show signs of moles' presence.

APPLICATION DIRECTIONS: Apply MOLE-MED with a hand-held sprayer or sprinkling can to entire area that is to be rid of moles or protected from moles. Cover treated area thoroughly with mixture of MOLE-MED and water. After treatment, water treated area with hose or sprinkler for an additional 25 minutes. Mole activity increases temporarily as moles leave the treated area. Cover the area to be treated thoroughly.

STORAGE AND DISPOSAL: Do not contaminate water, food or feed by storage or disposal.

STORAGE: Keep pesticide in original container. Do not put concentrate or dilute into food or drink containers. Store in a cool, dry place, preferably stored in a locked storage area.

PESTICIDE DISPOSAL: Wastes resulting from the use of this product may be disposed of on alte or at an approved waste disposal facility.

CONTAINER DISPOSAL: Triple rinse (or equivalent) then offer for recycling or reconditioning or puncture and dispose of in a sanitary landfill, or incineration, or if allowed by burning. If burned, stay out of smoke.

NOTICE: Buyer assumes all responsibility for safety and use not in accordance with directions.