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US ENVIRONMENTAL PROTECTION AGENCY
OFFICE OF PESTICIDES PROGRAMS
REGISTRATION DIVISION (TS-767)
WASHINGTON, DC 20460

NOTICE OF PESTICIDE: ☐ REGISTRATION ☒ REREGISTRATION

(Under the Federal Insecticide, Fungicide, and Rodenticide Act, as amended)

EPA REGISTRATION NO.	DATE OF ISSUANCE
655-19	SEP 29 1992

TERM OF ISSUANCE

NAME OF PESTICIDE PRODUCT

Prentox Warfarin Concentrate
Rax Powder

NAME AND ADDRESS OF REGISTRANT *(Include ZIP code)*

Prentiss Incorporated
C.B. 2000
21 Vernon St.
Floral Park, NY 11001

S426123 675 22

NOTE: Changes in labeling formula differing in substance from that accepted in connection with this registration must be submitted to and accepted by the Registration Division prior to use of the label in commerce. In any correspondence on this product always refer to the above U.S. EPA registration number.

On the basis of information furnished by the registrant, the above named pesticide is hereby Registered/Reregistered under the Federal Insecticide, Fungicide, and Rodenticide Act.

A copy of the labeling accepted in connection with this Registration/Reregistration is returned herewith.

Registration is in no way to be construed as an indorsement or approval of this product by this Agency. In order to protect health and the environment, the Administrator, on his motion, may at any time suspend or cancel the registration of a pesticide in accordance with the Act. The acceptance of any name in connection with the registration of a product under this Act is not to be construed as giving the registrant a right to exclusive use of the name or to its use if it has been covered by others.

Based on your response to the Reregistration Eligibility Document, EPA has reregistered the product listed above. Enclosed is a copy of your label stamped "Accepted". This action is taken under the authority of section 4(g)(2)(C) of the Federal Insecticide, Fungicide, and Rodenticide Act, as amended. Reregistration under this section does not eliminate the need for continual reassessment of pesticides. EPA may require submission of data at any time to maintain the registration of your product.

Make the revisions specified below before you print the final labeling for this product.

- In the third sentence under "CONTAINER DISPOSAL", insert a "," after "incineration" and after "authorities". The word "state" should be changed to "State".

Submit one copy of the final printed label before releasing the product for shipment with the revised labeling. Registrants of Warfarin products are allowed to sell or distribute products bearing old labeling until September 6, 1993.

Robert A. Forrest
Product Manager (14)
Registration Division (H7505C)

☐ ATTACHMENT IS APPLICABLE

SIGNATURE OF APPROVING OFFICIAL	DATE
	SEP 29 1992

EPA Form 8570-6 (Rev. 5-76) PREVIOUS EDITION MAY BE USED UNTIL SUPPLY IS EXHAUSTED. *U.S. GPO: 1988-312-018/80303

PRENTOX®

WARFARIN CONCENTRATE RAX® POWDER

For formulation only into registered end-use rodenticides to kill Norway rats, roof rats, and house mice.

ACTIVE INGREDIENT:	
Warfarin, 3-(alpha-Acetonylbenzyl)-4-hydroxycoumarin	0.5%
INERT INGREDIENTS:	99.5%
TOTAL:	100.0%

PRENTOX®—Registered Trademark of Prentiss Incorporated

KEEP OUT OF REACH OF CHILDREN

CAUTION

STATEMENT OF PRACTICAL TREATMENT

If swallowed: Call a Physician or Poison Control Center. Give a glass or two of water and induce vomiting by giving syrup of ipecac, if available, or by touching finger to back of throat. Do not give anything by mouth or induce vomiting if patient is unconscious.

If in eyes: Flush with plenty of water. Get medical attention if irritation persists.

If on skin: Wash off with soap and water

SEE SIDE PANEL FOR ADDITIONAL PRECAUTIONARY STATEMENTS

E.P.A. REG. NO. 655-19 **E.P.A. EST. NO. 12455 WI-1**

NET CONTENTS _____

LOT NO. _____

Manufactured by

PRENTISS INCORPORATED

Plant: Kaolin Road
Sandersville, GA. 31082
Office: C.B. 2000

PRECAUTIONARY STATEMENTS

HAZARDS TO HUMANS AND DOMESTIC ANIMALS

CAUTION

Keep away from humans, domestic animals and pets. Exposure during pregnancy should be avoided. Warfarin may cause harm to the fetus, including possible birth defects. If swallowed, this material may reduce the clotting ability of the blood and cause bleeding. When mixing baits, wear dust respirator and rubber gloves. If on skin, wash with soap and water.

NOTE TO PHYSICIAN: If ingested, administer Vitamin K1 intramuscularly or orally, as indicated in bishydroxycoumarin overdoses. Repeat as necessary based on monitoring of prothrombin times.

ENVIRONMENTAL HAZARD

This product is toxic to mammals and birds. Keep out of lakes, streams or ponds.

DIRECTIONS FOR USE

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

Only for formulation into a Rodenticide for (1) The following uses: control of Norway rats, roof rats, and house mice in and around homes, industrial buildings, and similar man-made structures; (2) Uses for which USEPA has accepted the required data and/or citations of data that the formulator has submitted in support of registration; and (3) Uses for experimental purposes that are in compliance with USEPA requirements.

Formulators are responsible for providing data to support their registration.

STORAG

Do not contaminate
or disposal.

STORAGE: Store in
not accessible to ch
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pesticide and wear
spills, wear protecti
prevent contact wi
Sweep onto a shove
salvage drum. Dispo

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