

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

EPA REGISTRATION NO.

655-19

DATE OF ISSUANCE

SEP 29 1992

NOTICE OF PESTICIDE: REGISTRATION

REREGISTRATION

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

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

5426123 675
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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 endorsement 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.

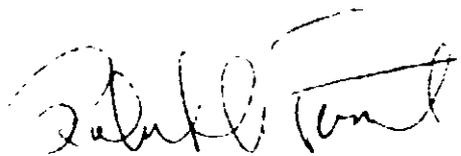
Based on your response to the Reregistration Eligibility Document, EPA has reregistered the product listed above. Enclosed is a copy of your label stamped "Accepted". This action is taken under the authority of section 4(g)(2)(C) of the Federal Insecticide, Fungicide, and Rodenticide Act, as amended. 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

PRENTOX^(R)

WARFARIN CONCENTRATE RAX[®] POWDER

**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.

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

ACTIVE INGREDIENT:

Warfarin, 3-(alpha-Acetylbenzyl)-
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**

STORAGE

Do not contaminate or disposal.

STORAGE: Store in not accessible to children or wildlife. If containing leaks. Take pesticide and wear protective gear. Sweep onto a salvage drum. Dispose

PESTICIDE DISPOSAL: Pesticide or Environmental Hazardous Waste Regional Office for

CONTAINER DISPOSAL: by shaking and tapping clinging particles. Then dispose at sanitary landfill, or to state and local authorities out of smoke.

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

