U.S. ENVIRONMENTAL PROTECTION AGENCY
Office of Pesticide Programs
Antimicrobials Division (7510C)
401 "M" St., S.W.
Washington, D.C. 20460

| EPA Reg. Number: | Date of Issuance: |
|---|---|
| 72643-1 | AUG 29 2000 |

Term of Issuance: Conditional

Name of Pesticide Product: DiSorb™ Tube

NOTICE OF PESTICIDE:
_x_ Registration
____ Reregistration

(under FIFRA, as amended)

Name and Address of Registrant (include ZIP Code):

Bohle Medical Supplies DiSorb™ Tube
11324 17th Avenue Court, N.W.
Gig Harbor, WA 98332

**Note:** Changes in labeling 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 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.

Registration is in no way to be construed as an endorsement or recommendation of this product by the 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.

This product is conditionally registered in accordance with FIFRA sec. 3(c)(7)(A) provided that you:

1. Submit and/or cite all data required for registration/ reregistration of your product under FIFRA sec. 3(c)(5) when the Agency requires all registrants of similar products to submit such data; and submit acceptable responses required for reregistration of your product under FIFRA section 4.

2. Make the following label changes:

   a. Revise the EPA Registration Number to read, "EPA Reg. No. 72643-1".

   b. Delete the "Reorder Number" from the label.

3. Submit two copies of the revised final printed label for the record.

If these conditions are not complied with, the registration will be subject to cancellation in accordance with FIFRA sec. 6(e). Your release for shipment of the product constitutes acceptance of these conditions.

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

| Signature of Approving Official: | Date: |
|---|---|
| Robert S. Brennis, PM 32 | AUG 29 2000 |

EPA Form 8570-6

2 93

Reorder Number: DYNDDS1500CC or DYNDDS3000CC

ACCEPTED
WITH COMMENTS
In EPA Letter Dated:

AUG 29 2000

Under the Federal Insecticide, Fungicide, and Rodenticide Act as amended for the pesticide registered under EPA Reg. No. 72643-1

# DiSorb™ Tube

Liquid Medical Waste Treatment Product
Solidifies Liquid Laboratory, Human and Animal Wastes
For use with all suction canisters

**Active Ingredients:**

| | |
|---|---|
| Iodine* | 10.65% |
| Sodium dichloroisocyanurate dihydrate** | 5.00% |
| Other Ingredients | 84.35% |
| **Total** | 100.00% |

*From povidone iodine and potassium iodide
**Provides 2.8% available chlorine

EPA Reg. No. 72643-R

EPA Est. No.

## KEEP OUT OF REACH OF CHILDREN

## CAUTION

| FIRST AID | |
|---|---|
| Have the product container or label with you when calling a poison control center or doctor, or going for treatment. | |
| **If in eyes** | – Hold eye open and rinse slowly and gently with water for 15-20 minutes. Remove contact lenses, if present, after the first 5 minutes, then continue rinsing eye.<br>– Call a poison control center or doctor for treatment advice |
| **If swallowed** | – Call a poison control center or doctor immediately for treatment advice.<br>– Have a person sip a glass of water if able to swallow.<br>– Do not induce vomiting unless told to do so by a poison control center or doctor<br>– Do not give anything by mouth to an unconscious person. |

## PRECAUTIONARY STATEMENTS

## HAZARDS TO HUMANS AND DOMESTIC ANIMALS

## CAUTION

Causes moderate eye irritation. Avoid contact with eyes or clothing. Harmful if swallowed. 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.

- Keep contents of box dry at all times.
- Wear hand and eye protection when handling infectious waste.
- Place canister as usual and open accessory port.
- Use the appropriate size DiSorb™ Tube based on the size of the suction canister. Each tube in this box contains 45 g of DiSorb™, which solidifies up to 1500 cc of liquid waste. OR Each tube in this box contains 85 g DiSorb™, which solidifies up to 3000 cc of liquid waste.
- Remove a DiSorb™ Tube from the box and shake for 3 seconds to disperse the product.
- Insert DiSorb™ Tube into the suction canister and close accessory port. NOTE: DiSorb™ Tube will dissolve when in contact with liquid.
- Connect suction tubing and use the canister as directed.

## STORAGE AND DISPOSAL

**STORAGE:** Store in a cool, dry, well ventilated area.
**PESTICIDE DISPOSAL:** Do not contaminate water, food or feed by storage or disposal. Wastes resulting from the use of this product may be disposed of on site or at an approved waste disposal facility.

## LIMITED WARRANTY AND DISCLAIMER

This manufacturer warrants (a) that this product conforms to the chemical description on the label; (b) that this product is reasonably fit for the purposes set forth in the directions for use when it is used in accordance with such directions; and (c) that the directions, warnings and other statements of this label are based upon reasonable experts' evaluation of reasonable test of effectiveness. THE MANUFACTURER NEITHER MAKES NOR INTENDS, NOR DOES IT AUTHORIZE ANY AGENT OR REPRESENTATIVE TO MAKE ANY OTHER WARRANTIES, EXPRESSED OR IMPLIED, AND IT EXPRESSLY EXCLUDES AND DISCLAIMS ALL IMPLIED WARRANTIES OF MERCHANTABILITY AND FITNESS FOR PARTICULAR PURPOSE. This warranty does not extend to, and the Buyer shall be solely responsible for, any and all loss or damage which results from the use of this product in any manner which is consistent with the label directions or warning.

BUYER'S EXCLUSIVE REMEDY AND MANUFACTURER'S OR SELLERS EXCLUSIVE LIABILITY FOR ANY AND ALL CLAIMS, LOSSES, DAMAGES, OR INJURIES RESULTING FROM THE USE OR HANDLING OF THIS PRODUCT, WHETHER OR NOT SUCH LIABILITY IN TORT OR OTHERWISE, SHALL BE LIMITED AT THE MANUFACTURER'S OPTION, TO REPLACEMENT OF, OR THE REPAYMENT OF THE PURCHASE PRICE FOR, THE QUANTITY OF PRODUCT WITH RESPECT TO WHICH DAMAGES ARE CLAIMED IN NO EVENT SHALL MANUFACTURER OR SELLER BE LIABLE FOR SPECIAL, INDIRECT OR CONSEQUENTIAL DAMAGES RESULTING FOR THE USE OF HANDLING OF THE PRODUCT.

**BOHLE MEDICAL SUPPLIES PTY LTD.**
9 Orchard Road; Brookvale, NSW 2100, Australia

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