

WJC

IODINE DISINFECTANT

Iodine Cleaner, Deodorizer, Disinfectant and Sanitizer

**FOR USE ON FARM PREMISES, POULTRY
HOUSES AND FOOD PLANTS.**

ACTIVE INGREDIENTS:

alpha-(p-nonylphenyl)-omega-hydroxy-poly [arylethylene]-iodine complex	18.05%
(Providing 1.75% available iodine)	
Phosphoric acid	18.00%
TOTAL ACTIVE INGREDIENTS	34.05%
INERT INGREDIENTS	65.95%
TOTAL INGREDIENTS	100.00%

DANGER KEEP OUT OF REACH OF CHILDREN

Practical Treatment: In case of contact, immediately flush eyes or skin with plenty of water for at least 15 minutes. For eyes, call a physician.

If swallowed, drink promptly a large quantity of milk, egg whites, gelatin solution or, if these are not available, drink large quantities of water. Call a physician.

NOTE TO PHYSICIAN: Excessive mucosal damage may contraindicate the use of gastric lavage. Measures against circulatory shock, respiratory depression, and convulsion may be needed.

For Medical, Industrial, Commercial, and Agricultural Use Only.

SOLD BY:

WADE JONES COMPANY, INC.
1100 Shaver Rd., Springdale, AR 72764

NET CONTENTS: 1 U.S. GAL. (3.79 l)

ACCEPTED
with COMMENTS
in EPA Letter Dated:

JUL 28 1987

Under the Federal Insecticide,
Fungicide, and Rodenticide Act
as amended, and the pesticide
laws of the State of EPA Reg. No.

48718-2

icide 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 section 3(c)(7)(A) provided that you:

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

2. Twelve months storage and stability data must be received by the Agency within 2 years after the granting of registration.

3. Insert a dash (-) between number "6" and the word "bis" in the second active ingredient.

4. Add the statement "May cause eye irritation." The If Swallowed and If On Skin statements should be deleted. Delete the statement "Harmful if Swallowed."

5. Add the phrase "EPA Registration No. 54614-3" to your label before you release the product for shipment.

6. Submit two (2) copies of your final printed labeling before you release the product for shipment. Refer to the A-79 Enclosure for further description of final printed labeling.

BEST AVAILABLE COPY

ATTACHMENT IS APPLICABLE

SIGNATURE OF APPROVING OFFICIAL *[Signature]* DATE

EPA Form 8570-6 (Rev. 5-76)

PREVIOUS EDITION MAY BE USED UNTIL SUPPLY IS EXHAUSTED.

92160: I: Pringle: P-1: KENCQ: 7/23/87: 8/3/87: cdb: vo: cdb

Pringle 7127