	PM 32 69687-1
	US ENVIRONMENTAL PROTECTION AGENCY OFFICE OF PESTICIDES PROGRAMS REGISTRATION NO. 69687-1 TERM OF ISSUANCE WASHINGTON, DC 20460
	NOTICE OF PESTICIDE: REGISTRATION NAME OF PESTICIDE PRODUCT
İ	(Under the Federal Insecticide, Fungicide, and Rodenticide Act, as amended) Super-Chlor
1	NAME AND ADDRESS OF REGISTRANT (Include ZIP code)
	Medtrol, Inc. 7157 N. Austin Avenue Niles, IL 60714
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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 the 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 un 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 prote health and the environment, the Administrator, on his motion, may at any time suspend or cancel the registration of a pelicide 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 cover by others.
	This product is conditionally registered in accordance with FIFRA section $3(c)(7)(A)$ provided that you:
	 Submit and/or cite all data required for registration/re- registration of your product under FIFRA section 3(c)(5) when the Agency requires all registrants of similiar products to submit such data; and submit acceptable responses required for reregistration of your product under FIFRA section 4.
	2. Make the labeling changes listed below before you release the product for shipment:
	a. Add the phrase "EPA Registration No. 69687-1"
	b. The ingredient statement term "Active Ingredients" should be singular.
	c. "Claim Super-Chlor contains 5.25% sodium hypochlorite solution diluted 1:10 with distilled water" is mis- leading and it should be revised.
	It may be similar to: Super-Chlor is 1:10 dilution of 5.25% sodium hypochlorite with distilled water.

3. Submit five (5) copies 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.

If these conditions are not complied with, the registration will be subject to cancellation in accordance with FIFRA section 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.

Sincerely yours,

Vivian A. Turner

Acting Product Manager (32) Antimicrobial Program Branch Registration Division (7505C)

Enclosure

Premoistened 8"x10" Towelette

CAT#SC110

SUPER-CHLORT

For wiping environmental surfaces & patient care equipment.

Active Ingredients: Sodium hypochlorite 0.525% *Inert Ingredients*: 99.475% *Inert Ingredients do not include weight of lowelette.

SUGGESTED AREAS OF USAGE:

- Medical, Dental and Laboratory Counters
- •Exam Tables, Carts, Point-of-Care Equipment
- •Telephones, Sink Tops, Toilet Seats

See panel on reverse for: Directions for Use

KEEP OUT OF REACH OF CHILDREN

- •If swallowed, drink large amounts of water. Do not induce vomiting.
- •If contact with eyes occurs, flush with water for 15 minutes. Get prompt medical attention.
- *If contact with skin occurs wash with plenty of soap and water,

FOR INSTITUTIONAL AND COMMERCIAL USE ONLY

*Do not reuse empty packet and towelette *Discard used towelette in designated waste container.



EPA Reg. No. _____ EPA Est #069687-IL-001

7157 Austin Avenue., Niles, IL 60714

PH: 800-647-7180

Nel Contents 0.6 fl. oz

Directions for use:

- -It is a violation of Federal Law to use this product in a manner inconsistent with its labeling
- -Always use personal protective equipments
- Open Super-Chlor Packet
- •Remove premoistened 8"x10" towelette
- ·Apply towelette and wipe desired surface to be decontaminated
- ·Allow to air dry and discard (see front panel)

This product is not to be used as a terminal sterilantifrigh level disinfectant on any surface instrument that (1) is introduced directly into the human body, either into or in contact with the bloodstream or normally sterile areas of the body, or (2) contacts intact mucous membranes but which does not ordinarilly penetrate the blood barriers or otherwise enter normally sterile areas of the body. This product may be used to precise nor decontaminate critical or semi-critical medical devices prior to sterilization or high level distribution.

Super-Chlor contains 5.25% Sodium hypochlorite solution diluted 1:10 with distilled water to comply with recommended infection control protocol.

Complete product line available by calling:

MED ROL

7157 Austin Avenue Niles, IL 60714 PH: 800-647-7180

ACCEPTED with COMMENTS in EPA Letter Dated:

OCT 24 1996

Under the Federal Insecticide, Fungicia, and Redemicide Act as amended, for the posticide registered under EPA Reg. No. 69687-/

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