

PM13

Reg. No. 11715-183

1/3

MAY 21 1992

11715-183

Until Reregistration

Flea Stop Yard Spray I

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Pet Chemicals Division of Shirlo, Inc.
4242 B.F. Goodrich Boulevard
Memphis, TN 38118

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

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

2. Make the labeling changes listed below before you release the product for shipment:

- a. Add the phrase "EPA Registration No. 11715-183."
- b. Under Precautionary Statements change "may cause eye irritation" to "causes moderate eye irritation."
- c. Under Statement of Practical Treatment add "Get medical attention" to the "If on Skin" statement.

63222:I:WP50:Moats:WP13-5:KEVRIC:05/20/92:06/19/92:DD

d. Revise the first sentence under Environmental Hazards to read:

This product is extremely toxic to fish and other aquatic organisms.

e. Revise the front panel statement to read: "For Use On Residential Lawns Only to Kill Fleas."

f. Under "Note" add "Keep children and pets off treated areas until spray has dried."

3. The following comments apply to chemistry for this product:

a. On the Confidential Statement of Formula (CSF), provide the pH of the product or the pH at a specified water dilution.

b. A one-year storage stability study must be conducted. The formulated product must be analyzed for its active ingredient at time zero and through a year of storage. The storage should be in warehouse conditions of temperature and humidity and stored in similar containers you will be using in the trade. Note: For the Storage Stability study you cannot reference the concentrate you are using to formulate your product.

Note: In reference to your letter dated June 15, 1990 you questioned if the acutes approved for EPA Registration No. 4758-157 can support this product. The answer is yes, since both products contain the same active and inert ingredients, the only difference is that the subject product contains only one-half of the amount contained in the registered product; except for the 4% increase in mineral spirits in the solvent which is rather unlikely to increase any hazard potential. Therefore, no further acute toxicity studies are required for the subject product.

4. 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.

George T. LaRocca
Product Manager (13)
Insecticide-Rodenticide Branch
Registration Division (H7505C)

Enclosures

