

**NITROFURAZONE SOLUBLE DRESSING- nitrofurazone ointment
VEDCO INCORPORATED**

EQUI-PHAR NITROFURAZONE SOLUBLE DRESSING

INDICATIONS:

For the prevention or treatment of surface bacterial infections of wounds, burns, and cutaneous ulcers.

ADMINISTRATION:

Apply directly on the lesion with a spatula, or first place on a piece of gauze. Application of a bandage is optional.

This preparation should be in contact with the lesion for at least 24 hours. The dressing may be changed several times daily or left on the lesion for a longer period.

PRECAUTION:

In case of deep or puncture wounds or serious burns, use only as recommended by a veterinarian. If redness, irritation, or swelling persists or increases, discontinue use, reconsult veterinarian. Avoid exposure to alkaline material and fluorescent lighting.

KEEP AWAY FROM EXCESSIVE HEAT OR DIRECT SUNLIGHT.

CONTENTS:

0.2% Nitrofurazone in a Water-soluble base of Polyethylene Glycols.

Do not use in horses intended for human consumption.

HUMAN WARNINGS:

CARCINOGENESIS: NITROFURAZONE, THE ACTIVE INGREDIENT OF NITROFURAZONE SOLUBLE DRESSING, HAS BEEN SHOWN TO PRODUCE MAMMARY TUMORS IN RATS AND OVARIAN TUMORS IN MICE.

SOME PEOPLE MAY BE HYPERSENSITIVE TO THIS PRODUCT. EITHER WEAR GLOVES WHEN APPLYING, OR WASH HANDS AFTERWARDS.

STORAGE:

Store at controlled room temperature between 15°-30°C (59°-86°F). Keep container tightly closed when not in use.

DISPLAY PANEL

For Use In Horses Only

An Antibacterial Preparation for Topical Application

CAUTION: FEDERAL LAW PROHIBITS THE USE OF THIS PRODUCT IN FOOD-PRODUCING ANIMALS.

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

Distributed By: VEDCO, INC., St. Joseph, MO 64507

Iss. 09-09

NET CONTENTS:

1 lb (453.6 g)

IMAGE OF LABEL

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<p>Iss. 09-09</p>	<p>ANADA 200-425, Approved by FDA</p>	<p>Lot No. _____ Exp. Date _____</p> <p style="text-align: center; color: blue; font-weight: bold;">No Varnish</p>

IMAGE OF LABEL

NITROFURAZONE SOLUBLE DRESSING			
nitrofurazone ointment			
Product Information			
Product Type	OTC ANIMAL DRUG LABEL	Item Code (Source)	NDC:50989-165
Route of Administration	TOPICAL	DEA Schedule	
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	NITROFURAZONE (NITROFURAZONE)	NITROFURAZONE	0.2 g in 1 g
Product Characteristics			
Color	yellow (YELLOW)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			
Packaging			
#	Item Code	Package Description	Marketing Start Date
1	NDC:50989-165-26	12 in 1 CASE	Marketing End Date

1	453.6 g in 1 JAR		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANADA	ANADA200425	11/14/2014	

Labeler - VEDCO INCORPORATED (021634266)

Revised: 11/2014

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