

**SCARLET OIL WOUND DRESSING- scarlet oil solution**  
**Durvet, Inc.**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**SCARLET OIL WOUND DRESSING**

**For Simple Wounds, Cuts and Abrasions**

Active Ingredient 0.3% (w/w) PCMX (Parachlorometaxlenol)

FOR ANIMAL USE ONLY

**INDICATIONS:**

For use as a dressing for simple wounds, cuts and abrasions on horses and mules.

**CAUTION:**

Use only as directed. Avoid contact with eyes and mucous membranes. Do not apply to large areas of broken skin. In case of deep or puncture wounds or serious burns, consult a veterinarian. If redness, irritation, or swelling persists or increases, discontinue use and consult a veterinarian.

**DIRECTIONS FOR USE:**

Clean the affected area, clipping hair if necessary. Direct spray at the site to be treated. Hold container 4 to 6 inches from animal. Apply freely as an open wound treatment or wrap with a clean bandage. Treatment may be applied once or twice daily.



**KEEP OUT OF REACH OF CHILDREN**

**INACTIVE INGREDIENTS:**

Mineral Oil, Isopropyl Alcohol, Methyl Salicylate, Benzyl Alcohol, Pine Oil, Eucalyptus Oil and Scarlet Red.

**WARNING: Not for use on animals intended for food.**

**STORAGE:**

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

**WARNING! FLAMMABLE! KEEP AWAY FROM HEAT AND OPEN FLAME.**

**Net Contents:**

16 OZ (473 mL)

Manufactured for:

**DURVET, INC.**

Blue Springs, Missouri 64014 Rev. 04-16

[www.durvet.com](http://www.durvet.com)

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**ISS21XB06**  
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**TAKE TIME**  
  
**OBSERVE LABEL DIRECTIONS**

NDC 30798-626-31  
  
**Scarlet Oil Wound Dressing**  
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Active Ingredient 0.3% (w/w) PCMX (Parachlorometaxyleneol)  
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Lot No. Exp. Date  
**NO VARNISH**



**SCARLET OIL WOUND DRESSING**

scarlet oil solution

**Product Information**

<b>Product Type</b>	OTC ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:30798-626
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>CHLOROXYLENOL</b> (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	2.5 mg in 1 mL

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30798-626-31	12 in 1 CASE		
1		473 mL in 1 BOTTLE		

  

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/06/2015	

**Labeler** - Durvet, Inc. (056387798)

### Establishment

Name	Address	ID/FEI	Business Operations
FIRST PRIORITY INCORPORATED		179925722	manufacture, label

Revised: 11/2022

Durvet, Inc.