

**HORSEMANS DREAM FUNG-A-WAY- benzalkonium chloride solution**  
**Manna Pro Products LLC**

*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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**CORONA FUNG-A-WAY TOPICAL FUNGICIDE SOLUTION**

Drug Facts

Active Ingredients: Benzalkonium chloride 0.15%

Purpose: Skin wound cleaner

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Uses:

For use on horses and dogs as an aid in the control of ringworm, summer itch, girth itch, and other fungal infections.

Warnings

For external veterinary use only.

Not for human use.

Not for use on animals intended for food

When using this product avoid eyes and mucous membranes. If contact occurs, flush immediately with water.

Stop use and consult your veterinarian if no improvement is noted within seven days.

Keep out of reach of children. In case of contact with eyes or mucous membranes, seek medical attention.

Directions

Soak affected area liberally with topical fungicide solution. Leave treated area uncovered.

Apply daily until hair begins to grow.

Rinse treated areas with clear water before reapplying.

Results should be apparent in a matter of days. Care should be taken to minimize the

animal's ingestion of topical fungicide solution through self-grooming.

Note: Efficacy is neutralized by soap or detergent residues.

Inactive Ingredients:

Allantoin, Benzethonium Chloride, Carbamide, FD&C Yellow #5, FD&C Red #40, Purified Water

CORONA

SINCE 1906

TOPICAL FUNGICIDE

FUNG-A-WAY

Aids in The Control Of Fungal Infections

- Topical Antiseptic to Help Control Ringworm, Summer Itch, Girt Itch, and Other Fungal Conditions.

- Protects and Soothes Irritated Skin

- Non-Staining

- For Use on Horses & Dogs

Distributed By:

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**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.15 g in 100 mL

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
<b>1</b>	NDC:86077-1040-6	453.6 mL in 1 BOTTLE		
<b>2</b>	NDC:86077-1040-2	907.2 mL in 1 BOTTLE		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
unapproved drug other		01/02/2017	

**Labeler** - Manna Pro Products LLC (147666812)**Registrant** - Manna Pro Products LLC (147666812)

Revised: 12/2025

Manna Pro Products LLC