

CHLORHEX 2X- chlorhexidine spray

Vedco, 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](#).

ChlorHex 2X

Indications:

For Mild to severe dermatological conditions where a full strength antiseptic, antimicrobial and deodorizing formulation may be of benefit.

Precautions:

For topical use only on Dogs, Cats and Horses. Avoid contact with eyes. If irritation develops, discontinue use and consult your veterinarian.

Storage

Store product at room temperature.

Active Ingredient:

Chlorhexidine Gluconate 4%

Inactive Ingredients:

Allantoin, Aloe Vera Gel, FD&C Blue #1, Fragrance, Isopropyl Alcohol, Polysorbate, Purified Water

Directions for Use:

Shake well before each use. Spray directly onto the affected area(s) 2-3 times daily, or as directed by veterinarian. Distract animal for several minutes following treatment to prevent licking.

Product Description

ChlorHex 2X 4% Spray is a maximum strength antimicrobial spray containing penetrating agents, allantoin and aloe vera for optimal therapeutic effectiveness.

Keep Out of Reach of Children

For Use on Dogs, Cats and Horses

For Animal Use Only

Package Label

Label for 8oz product:

NDC # 50989-918-24

ChlorHex 2X

4%

SPRAY

an antiseptic antimicrobial spray formulation
with aloe vera and allantoin

FOR USE ON DOGS, CATS & HORSES

For animal use only
Keep out of reach of children

VEDCO

Net Contents 8 FL. OZ. (240mL)

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



ChlorHex Spray

Distributed By

VEDCO, INC.
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Other Safety Information

TAKE TIME
OBSERVE LABEL DIRECTIONS

CHLORHEX 2X

chlorhexidine spray

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:50989-918
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	40.000 g in 1.0 L

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50989-918-24	0.240 L in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		10/11/2011	

Labeler - Vedco, Inc (021634266)**Registrant** - Continental Manufacturing Chemist, Inc. (005278007)

Revised: 6/2012

Vedco, Inc