EQUISHIELD CK HC- chlorhexidine, ketoconazole, hydrocortisone spray Creative Science

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.

EquiShield CK HC SPRAY

NDC 53413-027-06 - 16 oz NDC 53413-027-08 - 8 oz Drug Facts:

Active Ingredients:

 Chlorhexidine Gluconate
 2% (w/v)

 Ketoconazole
 1% (w/v)

 Hydrocortisone
 0.5% (w/v)

Purpose:

Antiseptic

Antifungal

Anti-Inflammatory

Inactive Ingredients:

Ethanol, Propylene Glycol, Purified Water, DMDM Hydantoin, Fragrance, FD&C Yellow #6

Indications:

For dermatological conditions responsive to Chlorhexidine and Ketoconazole. May be used for skin conditions with an inflammatory component. For use on horses, dogs and cats.

Warnings:

For external use only.

Avoid contact with eyes and mucous membranes.

Consult your veterinarian before use.

Keep out of reach of children.

Directions:

Apply directly to affected areas as directed by your veterinarian.

Other Information:

- Store at room temperature 15° 30°C (58° 86°F).
- Store in a cool, dry place.

REV0825 9007-03-00 UPC 851559000726 16 oz

REV0825 9007-03-01 UPC 851559000948 8 oz

Manufactured for Creative Science

1700 Albany Place SE, Orange City, IA 51041

Phone: 888-506-3039 www.creative.science



Equishield CK



Chlorhexidine Gluconate 2% Ketoconazole 1% Hydrocortisone 0.5%

Antiseptic, Anti-inflammatory, & Antifungal Spray

FOR VETERINARY USE ONLY KEEP OUT OF THE REACH OF CHILDREN

Net Contents: 8 fl oz (236 ml)

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Antisptic & Anti-inflammatory & Antifungal Spray

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EQUISHIELD CK HC

chlorhexidine, ketoconazole, hydrocortisone spray

Product Information

Product Type OTC ANIMAL DRUG Item Code (Source) NDC:53413-027

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE -**CHLORHEXIDINE** 20 mg UNII:R4KO0DY52L) **GLUCONATE** in 1 mL

10 mg HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYDROCORTISONE in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53413-027-08	236 mL in 1 BOTTLE, SPRAY		
2	NDC:53413-027-06	473 mL in 1 BOTTLE, SPRAY		

Marketing Information

Application Number or Monograph Marketing Citation Category

Marketing Start Date

Marketing End Date

unapproved drug other	08/01/2025	

Labeler - Creative Science (609777461)

Registrant - Creative Science (609777461)

Revised: 12/2025 Creative Science