

EQUISHIELD CK HC- chlorhexidine, ketoconazole, hydrocortisone spray
Kinetic Technologies, 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.

EquiShield CK HC SPRAY

Drug Facts:

Active Ingredients:

Chlorhexidine Gluconate 2% (w/v).....Antiseptic
Ketoconazole 1% (w/v).....Antifungal
Hydrocortisone 0.5% (w/v).....Anti-inflammatory

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.
Keep out of reach of children.
Consult your veterinarian before use.

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.

NDC 51031-027-08

NDC 51031-027-06

Made in U.S.A.

#9007-03-01

#9007-03-00

KINETIC™

P.O. Box 12388, Lexington, KY 40583

877.786.9882 • Fax: 859.258.9177

www.KineticVet.com

KINETICVET®

EquiShield® CK HC SPRAY

Chlorhexidine Gluconate 2%

Ketoconazole 1%

Hydrocortisone 0.5%

Antisptic & Anti-inflammatory Spray

FOR VETERINARY USE ONLY

KEEP OUT OF THE REACH OF CHILDREN

Net Contents: 8 fl oz (236 ml)

Net Contents: 16 fl oz (473 ml)



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HC SPRAY

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Antiseptic & Anti-inflammatory Spray

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EquiShield[®]CK HC NDC 51031-027-08
SPRAY

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Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:51031-027
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	20 mg in 1 mL
KETOCONAZOLE (UNII: R9400W927I) (KETOCONAZOLE - UNII:R9400W927I)	KETOCONAZOLE	10 mg in 1 mL
HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	5 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51031-027-08	236 mL in 1 BOTTLE, SPRAY		
2	NDC:51031-027-06	473 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/01/2023	

Labeler - Kinetic Technologies, LLC (164935731)

Registrant - Kinetic Technologies LLC (164935731)

Revised: 8/2023

Kinetic Technologies, LLC