

PROVITA KONQUEST- formic acid gel
Provita Eurotech Ltd

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.

Hoofsure Konquest

A proprietary blend of organic acids and essential oils with potent penetrating action. Does not contain copper or antibiotics.

Directions for Use

For use in sheep, cattle, horses and goats. Clean and dry hooves thoroughly before use. Pare hoof, if necessary. Apply the gel to the affected area using a standard caulking gun. Wipe end of the nozzle with a clean cloth and replace cap tightly after each use. Can be used with a hoof bandage. After a maximum of two days, remove the wrap. Where necessary, repeat application.

Precautions

Care must be taken when handling product.

Wear suitable protective clothing and eye protection.

Do not breathe vapour.

Contains Formic Acid.

Causes burns.

In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.

Keep out of reach of children.

See Health and Safety Data Sheet.

For external use only.

For animal use only.

Keep Cool.

Ingredients

Organic acids (Acetic, Formic and Salicylic), Pyrrolidone, Water, Hydroxyethylcellulose, Tea-tree Oil, Ethanol, Alkoxylated Cetyl Alcohol, Laureth-4, E124, E133

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PRINCIPAL DISPLAY PANEL - 300 g Cartridge Label

Provita

Animal Health Naturally

ANTIBIOTIC

FREE

Hoofsure

Konquest

Topical hoof gel

30

Applications

300g

**ANTIBIOTIC
FREE**

Hoofsure Konquest

Topical hoof gel



A proprietary blend of organic acids and essential oils with potent penetrating action. Does not contain copper or antibiotics.

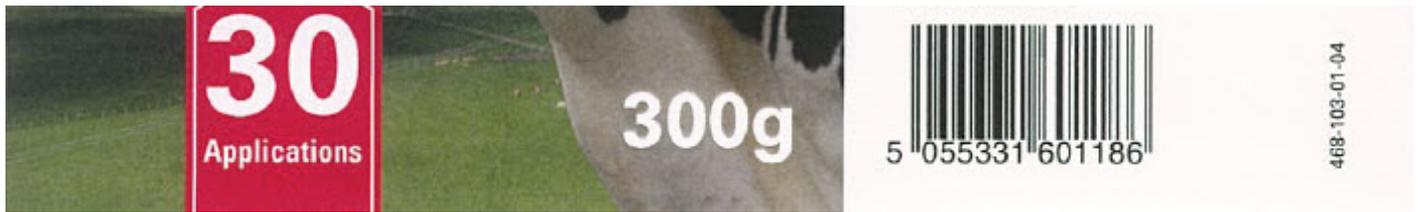
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Danger: Flammable liquid and vapour. Harmful if swallowed. Causes severe skin burns and eye damage. Keep away from heat/ sparks/open flames/hot surfaces — No smoking. Toxic if inhaled. May damage the unborn child. Care must be taken when handling product. Do not breathe dust / fume / gas / mist / vapours / spray. Wear protective gloves/protective clothing/ eye protection/face protection. IF SWALLOWED: Rinse mouth. Do NOT induce vomiting. IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing.

Rinse skin with water/ shower. IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do, continue rinsing. If symptoms persist, immediately call a POISON CENTER or doctor/physician. In case of fire: Use water, carbon dioxide (CO₂), foam, dry powder for extinction. Store in a well-ventilated place. Keep cool. Keep out of reach of children. For animal use only.



PROVITA KONQUEST

formic acid gel

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:16371-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FORMIC ACID (UNII: 0YIW783RG1) (FORMIC ACID - UNII:0YIW783RG1)	FORMIC ACID	14.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
LAURETH-4 (UNII: 6HQ855798J)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
PONCEAU 4R (UNII: Z525CBK9PG)	
ACETIC ACID (UNII: Q40Q9N063P)	
TEA TREE OIL (UNII: VIF565UC2G)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
PYRROLIDONE (UNII: KKL5D39EOL)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	

Product Characteristics

Color	BLUE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16371-104-02	48 in 1 CARTON		
1	NDC:16371-104-01	300 g in 1 CARTRIDGE		
2	NDC:16371-104-04	12 in 1 CARTON		

2	NDC:16371-104-03	15 g in 1 SYRINGE, PLASTIC		
3	NDC:16371-104-06	12 in 1 CARTON		
3	NDC:16371-104-05	30 g in 1 SYRINGE, PLASTIC		
4	NDC:16371-104-08	48 in 1 CARTON		
4	NDC:16371-104-07	300 g in 1 JAR		
5	NDC:16371-104-09	72 in 1 CARTON		
5	NDC:16371-104-07	300 g in 1 JAR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		01/01/2012	

Labeler - Provita Eurotech Ltd (238318232)

Establishment

Name	Address	ID/FEI	Business Operations
Provita Eurotech Ltd		220688192	API MANUFACTURE, MANUFACTURE, PACK, LABEL, ANALYSIS

Revised: 8/2023

Provita Eurotech Ltd