

DERMOPLAST FIRST AID- benzethonium chloride and benzocaine spray
Advantice Health, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Dermoplast® First Aid

Drug Facts

Active ingredients	Purposes
Benzethonium chloride 0.2%	First aid antiseptic
Benzocaine 20%	Topical analgesic

Uses

first aid for the temporary relief of pain and itching and to help prevent infection in minor cuts, scrapes and burns

Warnings

For external use only

Flammable do not use near heat, flame, or fire or while smoking

Allergy alert: do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine or other "caine" anesthetics.

Do not use

- in the eyes
- over large areas of the body

When using this product

- avoid contact with eyes. Do not spray in the face or mouth.
- use only as directed
- intentional misuse by deliberately concentrating or inhaling the contents can be harmful or fatal
- do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120°F.

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- itching, rash or irritation develops

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- clean the affected area

Adults and children 2 years of age and older	apply a small amount to affected area 1 to 3 times daily
Children under 2 years of age	consult a doctor

- to use this product, hold the can 6 to 12 inches away from the affected area. Direct spray nozzle towards skin and press button to activate spray.
- to apply to face, spray in palm of hand and gently apply

Other information

- avoid contact with leather, fabric and upholstery to prevent possible staining or discoloration
- store at 20-25°C (68-77° F)

Inactive ingredients

aloe vera gel (decolorized), cyclomethicone, dipropylene glycol, isobutane, SD alcohol 40-B (73% v/v), tocopheryl acetate (vitamin E acetate)

Questions?

1-800-345-0032

Mon - Fri 8AM- 5PM EST

Dermoplast.com

Distributed by Advantice Health, LLC
Cedar Knolls, NJ 07927

PRINCIPAL DISPLAY PANEL - 78 g Can Label

Dermoplast®

Antiseptic & Pain

Relieving Spray

HOSPITAL STRENGTH

FIRST AID

ANTIBACTERIAL SPRAY

KILLS 99% OF GERMS

To Prevent Infection

For Minor Cuts, Scrapes & Burns

Disinfects

Burns &

Wounds

Provides

Fast

Pain Relief

Soothing

Aloe &

Vitamin E

NET WT. 2.75 oz (78 g)

Drug Facts (continued)

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 Cedar Knolls, NJ 07927
 DOT 2P ☞ M5735 81-6701-03

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TYPE LIMIT

Dermoplast®

Antiseptic & Pain Relieving Spray

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FIRST AID
ANTIBACTERIAL SPRAY

KILLS 99% OF GERMS
To Prevent Infection

For Minor Cuts, Scrapes & Burns

+ Disinfects Burns & Wounds	🎯 Provides Fast Pain Relief	💧 Soothing Aloe & Vitamin E
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DERMOPLAST FIRST AID			
benzethonium chloride and benzocaine spray			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:16864-670
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	2 mg in 1 g	
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 g	
Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			
DIPROPYLENE GLYCOL (UNII: E107L85C40)			
CYCLOMETHICONE (UNII: NMQ347994Z)			

ALOE VERA LEAF (UNII: ZY81Z83H0X)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

ISOBUTANE (UNII: BXR49TP611)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16864-670-01	78 g in 1 CAN; Type 0: Not a Combination Product	01/01/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	01/01/2010	

Labeler - Advantice Health, LLC (192527062)

Revised: 3/2020

Advantice Health, LLC