

DERMOPLAST KIDS 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® Kids
First Aid Spray

Drug Facts

Active ingredients (To Deliver)	Purpose
Benzethonium chloride 0.2%	First aid antiseptic
Benzocaine 15%	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

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 barbadensis leaf juice, dimethyl ether, methylal, PEG-8

Questions?

1-800-345-0032

Mon - Fri 8AM - 5PM EST

Dermoplast.com

Distributed by
Advantice Health, LLC
Cedar Knolls, NJ 07927

PRINCIPAL DISPLAY PANEL - 57 g Can Label

Dermoplast®
KIDS

Antiseptic & Pain
Relieving Spray

FIRST AID
 ANTIBACTERIAL & PAIN
 RELIEVING SPRAY

STING FREE FORMULA

KILLS 99% OF GERMS
 To Help Prevent Infection†

For Minor Cuts, Scrapes & Burns

Disinfects Minor
 Burns & Wounds

#1
 Doctor
 Recommended
 Brand*

Provides Fast
 Pain Relief

NET WT. 2 oz (57 g)

<p>Drug Facts (continued)</p> <p>Directions • clean the affected area</p> <table border="1"> <tr> <td>Adults and children 2 years of age and older</td> <td>apply a small amount to affected area 1 to 3 times daily</td> </tr> <tr> <td>Children under 2 years of age</td> <td>consult a doctor</td> </tr> </table> <ul style="list-style-type: none"> • 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 <p>Other information • avoid contact with leather, fabric and upholstery to prevent possible staining or discoloration</p> <ul style="list-style-type: none"> • store at 20-25°C (68-77°F) <p>Inactive ingredients aloe barbadensis leaf juice, dimethyl ether, methylal, PEG-8</p> <p>Questions? 1-800-345-0032 Mon - Fri 8AM - 5PM EST Dermoplast.com</p>	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	 <p>Antiseptic & Pain Relieving Spray</p> <p>FIRST AID ANTIBACTERIAL & PAIN RELIEVING SPRAY</p>  <p>STING FREE FORMULA</p> <p>KILLS 99% OF GERMS To Help Prevent Infection†</p> <p>For Minor Cuts, Scrapes & Burns</p> <p>Disinfects Minor Burns & Wounds</p> <p>#1 Doctor Recommended Brand*</p> <p>Provides Fast Pain Relief</p> <p>NET WT. 2 oz (57 g)</p>	<p>Drug Facts</p> <table border="1"> <thead> <tr> <th>Active ingredients (To Deliver)</th> <th>Purpose</th> </tr> </thead> <tbody> <tr> <td>Benzethonium chloride 0.2%.....</td> <td>First aid antiseptic</td> </tr> <tr> <td>Benzocaine 15%.....</td> <td>Topical analgesic</td> </tr> </tbody> </table> <p>Uses first aid for the temporary relief of pain and itching and to help prevent infection in minor cuts, scrapes and burns</p> <p>Warnings For external use only</p> <p>Flammable do not use near heat, flame, or fire or while smoking</p> <p>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.</p> <p>Do not use • in the eyes • over large areas of the body</p> <p>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.</p> <p>Ask a doctor before use if you have • deep or puncture wounds • animal bites • serious burns</p> <p>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</p> <p>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. ▶</p>	Active ingredients (To Deliver)	Purpose	Benzethonium chloride 0.2%.....	First aid antiseptic	Benzocaine 15%.....	Topical analgesic
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benzethonium chloride and benzocaine spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:16864-660
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	150 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
METHYLAL (UNII: 7H1M4G2NUE)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIMETHYL ETHER (UNII: AM13FS69BX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16864-660-01	57 g in 1 CAN; Type 0: Not a Combination Product	03/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/01/2022	

Labeler - Advantice Health, LLC. (192527062)

Revised: 1/2023

Advantice Health, LLC.