DERMOPLAST PAIN RELIEVING- benzocaine, menthol spray
Advantice Health

Dermoplast Pain relieving Spray

Drug Facts

Active ingredients

Benzocaine 20%

Menthol 0.5%

Purpose

Topical analgesic

Uses

for temporary relief of pain and itching associated with • sunburn • insect bites • minor cuts • minor burns • minor skin irritations

Warnings

For external use only

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

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

When using this product

- avoid contact with eyes. Do not spray in the face of 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.

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

Adults and children 2 years of age and older	apply to affected area not more than 3 to 4 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

acetylated lanoline alcohol, aloe vera gel (decolorized), butane, cetyl acetate, hydrofluorocarbon 152a, methylparaben, PEG-400 monolaurate, polysorbate 85

Questions?

Mon - Fri 8AM- 5PM EST **1-800-345-0032Dermoplast.com**

Package Labeling:

DERMOPLAST PAIN RELIEVING

benzocaine, menthol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:16864-690
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTANE (UNII: 6LV4FOR43R)	
CETYL ACETATE (UNII: 4Q43814HXS)	
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)	

METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 85 (UNII: A7F3N56197)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16864-690-01	78 g in 1 CAN; Type 0: Not a Combination Product	03/18/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/18/2022	

Labeler - Advantice Health (192527062)

Revised: 12/2023 Advantice Health