DUKAL STING RELIEF PAD- benzocaine swab
Dukal LLC

Drug Facts

Active Ingredients

Benzocaine, 6% w/v

Isopropyl Alcohol 60% w/v

Purpose

Topical Anesthetic

Antiseptic

Directions

Apply to affected area 3-4 times daily. For adults and children 2 years of age and older. Children under 2 years; consult physician.

Do Not Use

- In the eyes. If contact occurs, rinse thoroughly with water

Stop Use

If irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.

Caution Flammable

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Inactive Ingredient

Purified Water

D07110607 Rev5

Principal Display Panel - 0.4 mL Pouch Label

DUKAL™
CORPORATION

REF
856

NDC 65517-0005-1
STING RELIEF PAD

For temporary relief of pain and itching associated with minor burns, scrapes and insect bites.

For External Use Only

1 Pouch

DUKAL CORPORATION • (631) 656-3800
Ronkonkoma, New York • www.dukal.com

Made in China, Hecho en China, Fabriqué en Chine

DUKAL™ CORPORATION

REF 856

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For temporary relief of pain and itching associated with minor burns, scrapes and insect bites.

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LATEX 2

Made in China, Hecho en China, Fabriqué en Chine

Drug Facts

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Benzocaine, 6% w/v	Topical Anesthetic
Isopropyl Alcohol 60% w/v	Antiseptic

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

Inactive Ingredient Purified Water

D07110607 Rev5

DUKAL STING RELIEF PAD

benzocaine swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65517-0005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	60 mg in 1 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.6 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65517-0005-1	0.4 mL in 1 POUCH; Type 0: Not a Combination Product	06/01/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/01/2010	

Labeler - Dukal LLC (791014871)

Revised: 10/2023 Dukal LLC