

DUKAL STING RELIEF PAD- benzocaine swab

Dukal Corporation

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.

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



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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 not final	part348	06/01/2010	

Labeler - Dukat Corporation (791014871)