

STING-KILL- benzocaine and menthol solution
Randob Ltd.

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 USP 20%

Menthol USP 1%

Purpose

(for pain) Topical Anesthetic

(anti-itch) Antipruritic

Keep Out of Reach of Children

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

Uses

- Temporarily relieves pain and itching of bee stings, insect bites and minor skin irritations.

Warnings

- **For external use only**
- **Avoid contact with eyes**

Get medical help right away if

- You are sensitive to insect venom. You may develop a serious reaction when stung.

Stop use and ask a doctor if

- Condition worsens
- Symptoms last for more than 7 days

Directions

Children under 2 yrs.

- Do not use
- Consult doctor

Adults and children 2 yrs. and older

- Apply to affected area as needed but not more than 3 to 4 times a day.
- (1) Bee stings only, remove stinger before treatment.
- (2) Tear foil pack at notch, exposing pad.
- (3) Apply immediately to bite or sting area. Dispose of pad after use.

Inactive Ingredients

FD&C Blue #1, FD&C Yellow #5, Isopropyl Alcohol 15%, PEG-8, Water.

Package/Label Principal Display Panel

#1 CHOICE OF PHARMACISTS

2 in 1 FORMULA

STING-KILL®

Kills the Pain and Stops the Itch with MAXIMUM STRENGTH Benzocaine and Menthol

For fast relief from:

Bee Stings • Mosquito Bites

Insect Bites • Jellyfish

MADE IN THE USA

EXTERNAL ANESTHETIC WIPES

SANITARY • CONVENIENT

8 DISPOSABLE SATURATED WIPES



STING-KILL

benzocaine and menthol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52412-250
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 mL
MENTHOL (UNII: L7T10EP3A) (MENTHOL - UNII:L7T10EP3A)	MENTHOL	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PEG-8 LAURATE (UNII: 762O8IWA10)	
WATER (UNII: 059QF0K00R)	

Product Characteristics

Color	BLUE (Blue)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52412-250-20	8 in 1 BLISTER PACK	01/01/1965	
1		0.56 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:52412-250-21	8 in 1 CARTON	01/15/2017	
2		0.56 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/01/1965	

Labeler - Randob Ltd. (061995007)

Registrant - Randob Ltd. (061995007)

