

STINGX- benzocaine swab
CoreTex Products Inc

Cortex StingX
65753-350

Active Ingredients

Benzocaine 6%

Purpose

Topical Analgesic

Use

For temporary pain relief from insect bites and stings

Warnings

For external use only

Do Not Use

- in or near eyes
- over large areas of the body
- over raw or blistered areas

Stop use and ask a doctor if

- conditions worsen or persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children

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

Directions

Apply to affected area not more than 3 to 4 times daily, for adults and children 2 years of age or older.

Inactive Ingredients

allantoin, glycerin, SD alcohol 40, water (aqua)

Made in USA for CoreTex Products, Inc.

Erwin, TN 37650 / www.CoreTexProducts.com

(877)684-5774

Principal Display Panel

Sting X Pouch

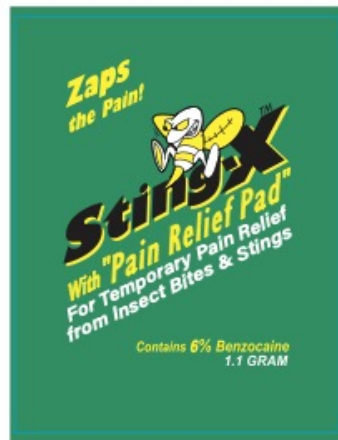
PMS 339 C



PMS 3945 C



PMS
Hexachrome
Black



Drug Facts	
Active Ingredients	Purpose
Benzocaine - 6%	Topical Anesthetic
Use: For temporary pain relief from insect bites and stings.	
Warnings	
For external use only	
Do not use - in or near eyes - over large areas of the body - over raw or blistered areas.	
Stop use and ask a doctor if conditions worsen or persist for more than 7 days or clear up and occur again within a few days.	
Keep out of reach of children: If swallowed get medical help or contact Poison Control Center right away.	
Directions: Apply to affected area not more than 3 to 4 times daily, for adults and children 2 years of age or older.	
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H2.5" x W2"

benzocaine swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65753-350
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	6 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
ALLANTOIN (UNII: 344S277G0Z)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65753-350-15	25 in 1 CONTAINER	11/25/2019	
1		1.1 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:65753-350-16	50 in 1 CONTAINER	11/25/2019	
2		1.1 mL in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:65753-350-17	300 in 1 CONTAINER	11/25/2019	
3		1.1 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 Drug	M017	11/25/2019	

Labeler - CoreTex Products Inc (061944620)

Establishment

Name	Address	ID/FEI	Business Operations
Cosmetic Enterprises		017701475	manufacture(65753-350)

Establishment

Name	Address	ID/FEI	Business Operations
CoreTex Products Inc		061944620	pack(65753-350)

Revised: 10/2023

CoreTex Products Inc