STINGX- benzocaine swab
CoreTex Products Inc

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.

Cortex StingX

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 aska 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

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

Made in USA for CoreTex Products, Inc.

Bakersfield, CA 93308

www.CoreTexProducts.com (877)684-5774

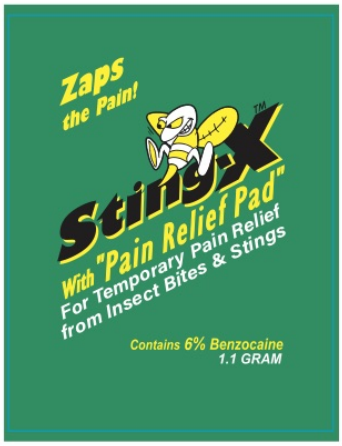
PRINCIPAL DISPLAY PANEL

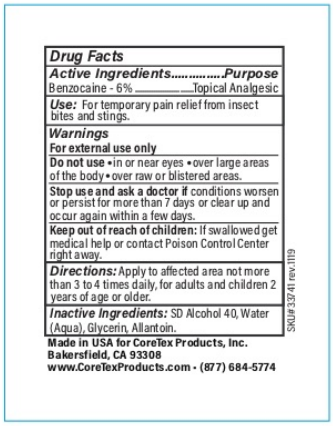
Sting X Pouch

PMS 339 C

PMS 3945 C

PMS Hexachrome Black





H2.5" x W2"

STINGX

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 not final	part348	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	label(65753-350)