BITE AND STING RELIEF WELL AT WALGREENS- benzocaine - 5.00% spray
Walgreens

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.

Drag Facts

Active ingredient	**Purpose**
Benzocaine - 5.00%	Pain Relief

Uses

For temporarily relief of pain and itching associated with insect bites

Warnings
For external use only

Flammable: do not use while smoking or near heat or flame

When using this product

- avoid contact with eyes
- use only as directed
- do not puncture or incinerate. Contents under pressure. Do not store at temperature above 120F

Stop use and ask a doctor if

• condition worsens
• symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of the children
In case of accidental ingestion, seek professional or contact a Poison Control Center immediately.

Direction

• shake well
• adults and children 2 years of age and older: Apply to affected area, not more than 3 to 4 times daily
• children under 2 years of age: Ask a doctor

Inactive ingredients

Aloe Barbadensis Leaf Juice, Ascorbic Acid, Camphor, Chamomilla Recutita (Matricaria) Flower Extract, Cholecalciferol, Diisopropyl Adipate,
Eugenia Caryophyllus (Clove) Flower Oil, Fragrance, Mentha Piperita (Peppermint) Oil, Octyldodecanol, Olea Europaea (Olive) Fruit Oil,
PEG-8 Dimethicone, Propylene Glycol, Pyridoxine HCl, Retinyl Palmitate, SD Alcohol 40, Silica, Sodium Propoxyhydroxypropyl Thiosulfate Silica,
Tocopheryl Acetate, Zea Mays (Corn) Oil

Questions and comments?

1-866-483-2846

BITE AND STING RELIEF WELL AT WALGREENS

benzocaine - 5.00% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-3190
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CHAMOMILE (UNII: FGL3685T2X)	
Cholecalciferol (UNII: 1C6V77QF41)	
Diisopropyl Adipate (UNII: P7E6YFV72X)	
CLOVE OIL (UNII: 578389D6D0)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
Octyldodecanol (UNII: 461N1O614Y)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
PEG-8 Dimethicone (UNII: GIA7T764OD)	
Propylene Glycol (UNII: 6DC9Q167V3)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
Alcohol (UNII: 3K9958V90M)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
Sodium Propoxyhydroxypropyl Thiosulfate Silica (UNII: 208G222332)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CORN OIL (UNII: 8470G57WFM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-3190-03	85 g in 1 CAN; Type 0: Not a Combination Product	03/15/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/15/2013	

Labeler - Walgreens (008965063)

Registrant - Product Quest Mfg (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(0363-3190), label(0363-3190)

Revised: 1/2018 Walgreens