AFTER BITE OUTDOOR- diphenhydramine hcl gel
Tender Corporation

After Bite Outdoor

Active Ingredient

Diphenhydramine HCl 2%

Purpose

Topical Analgesic

Uses

For the temporary relief of pain and itching associated with
- insect bites
- minor burns
- sunburn
- minor skin irritations
- minor cuts
- rashes due to poison ivy, poison oak, and poison sumac

Warnings

For external use only

Do not use

Do not use
- on large areas of the body
- with any other product containing diphenhydramine, even one taken by mouth

Ask a doctor before use

Ask a doctor before use
- on chicken pox
- on measles

When using

When using this product avoid contact with eyes.

Stop use and ask a doctor if

Stop use and ask a doctor if

- condition worsens
- if symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Directions

- do not use more than directed
- adults and children over 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 Vera, Citric Acid, Ethyl Alcohol, Glycerin, Methocel, Methylparaben, Oat Beta Glucan, Propylparaben, Purified Water, Sodium Hydroxide, Tea Tree Oil, Vitamin E

Package Labeling

AFTER BITE OUTDOOR

diphenhydramine hcl gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44224-0156
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ALCOHOL (UNII: 3K9958V90M)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
TEA TREE OIL (UNII: VIF565UC2G)	
METHYLCELLULOSE (100 CPS) (UNII: 4GFU244C4J)	
OAT (UNII: Z6J799EAJK)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-0156-1	1 in 1 BOX	03/01/2018	09/30/2020
1		20 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:44224-0156-0	20 g in 1 TUBE; Type 0: Not a Combination Product	03/01/2018	09/30/2020

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/01/2018	12/31/2024

Labeler - Tender Corporation (064437304)

Registrant - Tender Corporation (064437304)

Establishment

Name	Address	ID/FEI	Business Operations
Tender Corporation		064437304	manufacture(44224-0156)