

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 is a powerful itch treatment that provides immediate relief from insect bites and stings. This itch erasing gel contains Antihistamine and Tea Tree Oil to help soothe the skin and stop the pain and discomfort.

Perfect to carry on any outdoor adventure!

Drug Facts

Active ingredient	Purpose
Diphenhydramine HCl 2%	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 on large areas of the body with any other product containing diphenhydramine, even one taken by mouth

Ask a doctor before use on chickenpox on measles

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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

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Directions

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Inactive ingredients

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See our complete line of products at AfterBite.com

Insectlopedia.com Your source for treatment, prevention, and the latest buzz on bugs.

SATISFACTION GUARANTEED:
Return product to place of purchase or manufacturer.



NDC 44224-0156-1

After Bite®

After Bite®

After Bite®

OUTDOOR

After Bite®

**FAST ACTING
GEL FORMULA
with Antihistamine**

**POWERFUL
INSTANT ITCH
RELIEF!**

BE READY for



Mosquitoes, Bees & Wasps,
Biting Flies, and Other Insects

Net Weight: 0.7oz. (20g)

BE READY for



Mosquitoes



Biting Flies



Bees & Wasps



Other Insects

BE READY for



Mosquitoes



Biting Flies



Bees & Wasps



Other Insects

ART-0002-1211-1



After Bite® OUTDOOR

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: Z6J799EAKJ)	
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)

