# THE ITCH ERASER- diphenhydramine hydrochloride spray Adventure Ready Brands

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.

-----

#### The Itch Eraser Spray

#### **Drug Facts**

#### Active ingredients:

Diphenhydramine HCI 2%

#### Purpose:

Topical analgesic

#### Uses

### For temporary relief of pain and itching associated with:

- minor burns
- sunburn
- minor cuts
- insect bites
- minor skin irritations
- 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 chicken pox • on measles

#### When using this product

avoid contact with the eyes.

#### 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 children.

If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

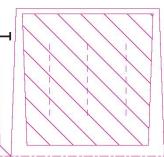
- do not use more often than directed
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily.
- children under 2 years of age: ask a doctor.

#### **Inactive Ingredients**

Glycerin, Oat Beta Glucan, Povidone, Purified Water, SD Alcohol 40-2, Sodium Bicarbonate (Baking Soda), Tea Tree Oil, Vitamin E Acetate, Zinc Acetate

### **Package Labeling**







MAX STRENGTH

**FAST ITCH** & PAIN RELIEF

**MOISTURIZES & NOURISHES** 

**RELIEF With:** 

STEROID-FREE



MAX STRENGTH

## ANTI SPRA

with Antihistamine Diphenhydramine HCI 2%

STOP THE ITCH



4.4 cm

WITH ANTIHISTAMINE

MAX STRENGTH

**TEMPORARY RELIEF OF** PAIN AND ITCHING ASSOCIATED WITH:



Poison Ivy, Oak, & Sumac



Minor Skin Irritations



Sunburn



The Itch Eraser® spray features a steroid-free antihistamine formula. This soothing spray provides fast itch and pain relief.

#### Drug Facts

Active ingredient
Diphenhydramine HCI 2%

Purpose Topical Analges

Uses
For the temporary relief of pain and litching associated with ■ minor burns ■ sunburn ■ minor cuts ■ insect bites ■ minor skin irritations ■ raises due to poison ivy, poison oak, and praison signat.

ooison 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 chicken pox ■ on measles

When using this product avoid contact with the

#### Stop use and ask a doctor if

a 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. If swallowed, get medical help or contact a Poison Control Center right away.

#### Directions

■ Shake before use
■ Remove safety tab from spray head
■ Adults and children 2 years of age and older:
apply to affected area not more than 3 to 4 times

■ Children under 2 years of age: consult a doctor

Inactive ingredients Glycerin, Oat Beta Glucan, Povidone, Purified Water, SD Alcohol 40-2, Sodium Bicarbonate (Baking Soda), Tea Tree Oil, Vitamin E Acetate, Tine Acetate.

From the makers of After Bite



#### THE ITCH ERASER

diphenhydramine hydrochloride spray

#### **Product Information**

HUMAN OTC DRUG NDC:90107-2410 **Product Type Item Code (Source)** 

**Route of Administration TOPICAL** 

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
- 11 - 11 - 11 - 11 - 11 - 11 - 11 -	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
OAT (UNII: Z6J799EAJK)		
ALCOHOL (UNII: 3K9958V90M)		
WATER (UNII: 059QF0KO0R)		
POVIDONE (UNII: FZ 989GH94E)		
GLYCERIN (UNII: PDC6A3C0OX)		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
TEA TREE OIL (UNII: VIF565UC2G)		
ZINC ACETATE (UNII: FM5526K07A)		

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:90107- 2410-1	1 in 1 BOX	09/01/2020		
1		28 mL in 1 BOTTLE, SPRAY; 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	09/01/2020		
final	pai to40	09/01/2020		

# Labeler - Adventure Ready Brands (064437304)

# **Registrant -** Adventure Ready Brands (064437304)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
Adventure Ready Brands		064437304	manufacture(90107-2410)		

Revised: 1/2023 Adventure Ready Brands