

ITCH ERASER- diphenhydramine hcl spray
Tender Corporation d/b/a Adventure Ready Brands

Itch Eraser Spray

Active Ingredient

Diphenhydramine HCl 2%

Purpose

Topical Analgesic

Uses

For the temporary relief of pain and itching associated with, minor burns, sunburn, minor cuts, insects 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 ox, measles

when using this product

avoid contact with eyes.

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

if swallowed, get medical help or contact poison control center right away.

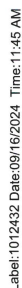
Directions

Shake before use

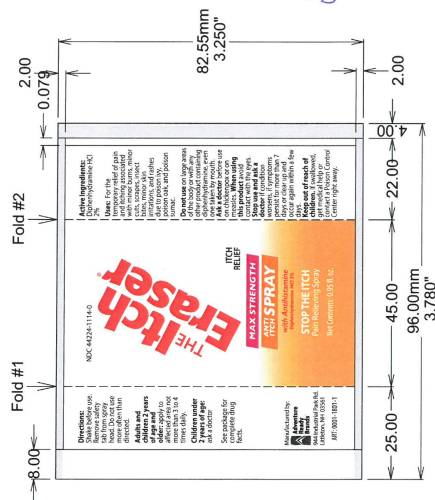
children under 2 years of age: consult a doctor

glycerinm oat beta glucanm povidone, purified water, sd alcohol 40-2, sodium bicarbonate (baking soda), tea tree oil, vitamine e acetate, zinc acetate

Itch Eraser Spray Label



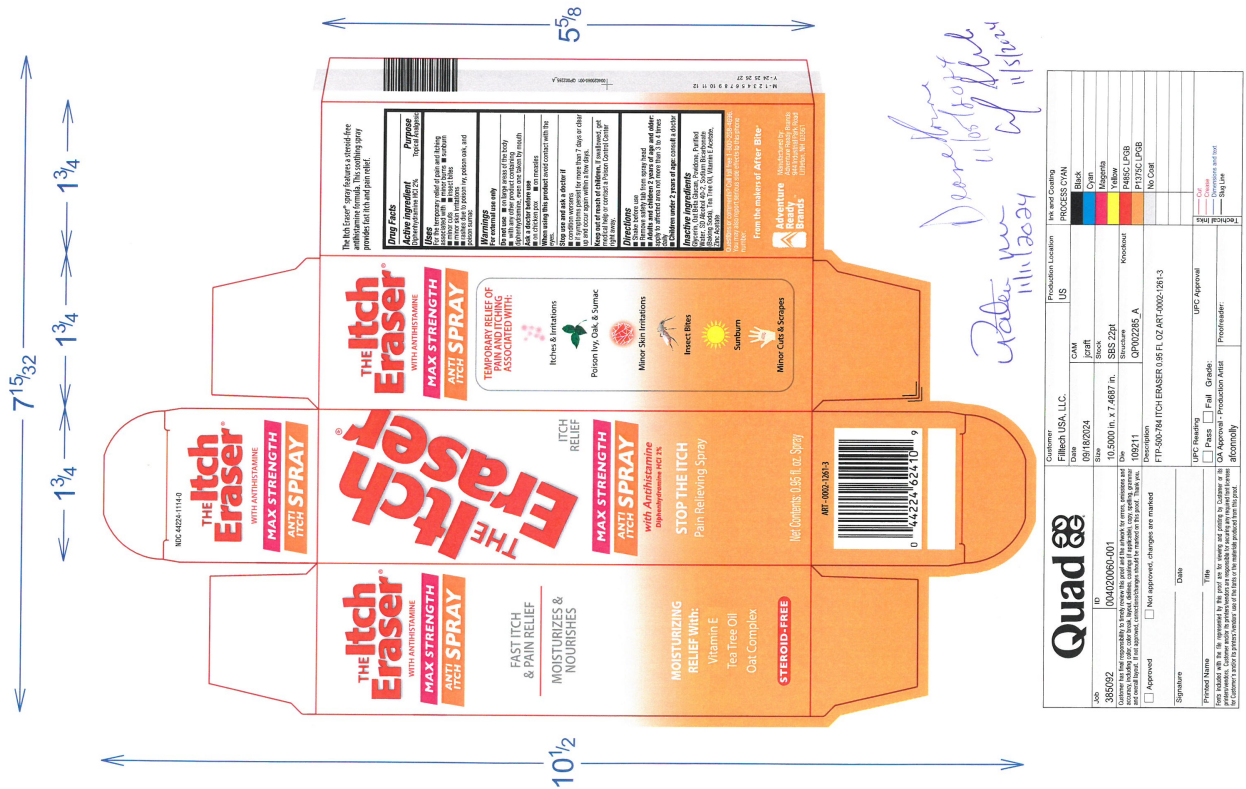
Metallic ink representation:	
Substrate representation:	
Adhesive representation:	
Varnish representation:	
White ink representation:	
Special representation:	



Done in
11/12/2024
of the
11/5/2024

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Itch Eraser Spray Box



ITCH ERASER

diphenhydramine hcl spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44224-1114
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII: 8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	0.51 g in 25.5 g

Inactive Ingredients

Ingredient Name	Strength
TEA TREE OIL (UNII: VIF565UC2G)	0.128 g in 25.5 g
POVIDONE K30 (UNII: U725QWY32X)	0.77 g in 25.5 g
WATER (UNII: 059QF0KO0R)	4.05 g in 25.5 g
OAT (UNII: Z6J799EAJK)	0.51 g in 25.5 g
ZINC ACETATE (UNII: FM5526K07A)	0.026 g in 25.5 g
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	0.128 g in 25.5 g
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.128 g in 25.5 g

GLYCERIN (UNII: PDC6A3C0OX)	0.77 g in 25.5 g
ALCOHOL (UNII: 3K9958V90M)	18.48 g in 25.5 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44224-1114-1	1 in 1 BOX	11/11/2024	
1	NDC:44224-1114-0	25.5 g 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 Drug	M017	11/11/2024	

Labeler - Tender Corporation d/b/a Adventure Ready Brands (064437304)

Establishment

Name	Address	ID/FEI	Business Operations
Tender corporation d/b/a Adventure Ready Brands		064437304	label(44224-1114) , pack(44224-1114)

Establishment

Name	Address	ID/FEI	Business Operations
Filltech USA, LLC		926433855	manufacture(44224-1114)