

ITCH RELIEF- diphenhydramine hcl and zinc acetate spray
FSA Store Inc.

Carring Mill Itch Relief Pump Spray

Active ingredients

Diphenhydramine HCl 2.0%,

Purpose

External analgesic

Uses

for the temporary relief of pain and itching associated with:

- minor burns
- minor cuts
- scrapes
- insect bites
- minor skin irritations

Warnings

For external use only.

Flammable:

Keep away from fire or flame.

Do not use

- on large areas of the body
- on chicken pox, poison ivy, or sunburn
- on broken, blistered, or oozing skin
- more often than directed
- with any other product containing diphenhydramine, even one taken by mouth

When using this product

- avoid contact with the eyes

Stop use and ask doctor if

- conditions worsens
- 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

- adults and children 2 years of age and older: spray on affected area not more than 3 to 4 times daily.
- children under 2 years of age: do not use, consult a doctor.

Other information

- store between 20° to 25°C (68° to 77°F)
- shake well before use
- pump must be primed before use

Inactive ingredients

SD ALCOHOL 40-B, WATER, GLYCERIN, CAMPHOR, TROMETHAMINE, PVP, ALOE BARBADENSIS LEAF JUICE

Questions?

1-866-964-0939

Principal Display Panel

caring mill

Extra Strength

Itch Relief

Pump Spray

Diphenhydramine Hydrochloride 2%/

External Analgesic

Easy-To-Spray Pump

- Relieves pain and itching associated

with minor burns, cuts, scrapes,

insect bites, and skin irritations

- Helps cool & soothe skin
- Ozone-Friendly, Non-Aerosol

2.5 FL. OZ. (74 mL)

INSIDE FACE

WIND #3

WIND #4

Front Label

Back Label (Peel & Reveal)

Drug Facts (continued)

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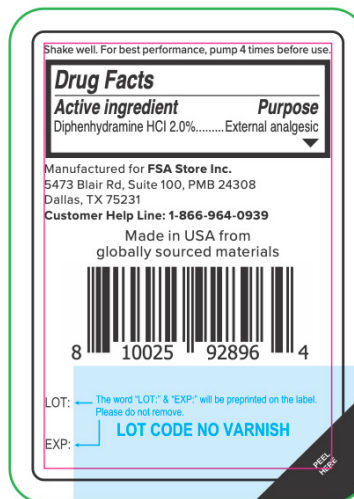
HINGE

1/16 round

2.4375 in



1.6875 in



FACE



BASE

ITCH RELIEF

diphenhydramine hcl and zinc acetate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81522-726
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 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAMPHOR (NATURAL) (UNII: N20HL7Q941)	
TROMETHAMINE (UNII: 023C2WHX2V)	
PVP (UNII: FZ989GH94E)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81522-726-01	74 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/19/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/19/2025	

Labeler - FSA Store Inc. (049283340)