ITCH RELIEF- itch relief gel gel
Neilmed Pharmaceuticals Inc.

Itch Relief

Drug Facts: Active Ingredients

Diphenhydramine HCl 2%

Purpose

Topical analgesic

Uses

Temporarily relieves pain and itching associated with:

- insect bites
- minor burns
- sunburn
- minor cuts
- scrapes
- 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

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- condition 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, 1-800-222-

Directions

- Do not use more 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

Other Information

Store at Room Temperature

Inactive ingredients

- Camphor
- diazolidinyl urea
- glycerin
- hydroxyethyl cellulose
- methylparaben
- propylene glycol
- propylparaben
- purified water
- SD alcohol 40-B
- sodium citrate

Tamper Evident

Do not use if seal on tube is punctured or missing

Questions?

1(877) 477-8633

Principal Display Panel

NM9999-Rev1-ItchRelief-3.5oz-TUBE-v4 Artwork Dielines

ITCH RELIEF

itch relief gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13709-351
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**
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
HYDROXYETHYL CELLULOSE (140 CPS AT 5%) (UNII: 8136Y38GY5)	
WATER (UNII: 059QF0KO0R)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13709-351-01	1 in 1 CARTON	11/26/2025	
1		103.5 mL in 1 TUBE; 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	11/26/2025	

Labeler - Neilmed Pharmaceuticals Inc. (799295915)

Establishment

Name	Address	ID/FEI	Business Operations
Neilmed Pharmaceuticals Inc.		799295915	manufacture(13709-351)

Revised: 11/2025 Neilmed Pharmaceuticals Inc.