A HEALTH ITCH RELIEF GEL- itch relief gel gel
Pharma Nobis, LLC

Freskaro Itch Relief Gel

Diphenhydramine HCl 2%

External Analgesic

Temporarily relieves pain due to: Minor burns, insect bites, sunburn, minor skin irritation, minor cuts, scrapes, rashes due to poison ivy, poison oak and poison sumac.

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
- measles

When using this product

- Avoid contact with eyes

Stop use and ask a doctor if

- condition gets worse
- symptoms last for more than 7 days
- symptoms clear up and occur again in a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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: do not use, consult a doctor.

Camphor, Citric Acid, Diazolidnyl Urea, Glycerin, Hydroxypropyl Methylcellulose, Methylparaben, Propylene Glycol, Propylparaben, SD Alcohol 38-B, Sodium Citrate, Purified Water.

Questions & comments?

1-833-551-0932

2026 Label

A HEALTH ITCH RELIEF GEL

itch relief gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82645-116
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
METHYLPARABEN (UNII: A2I8C7HI9T)	

PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82645-116-94	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/15/2026	

Labeler - Pharma Nobis, LLC (118564114)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(82645-116) , label(82645-116) , pack(82645-116) , analysis(82645-116)

Revised: 4/2026 Pharma Nobis, LLC