A HEALTH ITCH RELIEF GEL- itch relief gel gel
Bionpharma, Inc.

A+ Health Itch Relief Gel

Active Ingredient

Diphenhydramine HCl 2%

Purpose

External analgesic

Uses

Temporarily relieves pain due to: • minor burns • insect bites

• sunburn • minor skin irritation • minor cuts • scrapes • rashes

due to poison ivy, poison oak & 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 • 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.

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

Camphor, Citric Acid, Diazolidnyl Urea,

Glycerin, Hydroxypropyl Methylcellulose, Methylparaben, Propylene

Glycol, Propylparaben, SD Alcohol 38-B, Sodium Citrate, Purified Water.

Questions and comments?

call toll free 1-888-235-2466 (Mon - Fri 9AM - 5PM EST)

DISTRIBUTED BY: Bionpharma Inc., Princeton, NJ 08540

L0000940 R0125

†This product is not manufactured or distributed by the owner of the registered trademark Benadryl†®

4 fl oz (118 mL) tube

† Compare to the active ingredient

in **Benadryl ® Gel**

a+ health ™

extra strength

itch

relief

gel

diphenhydramine HCl 2%

topical analgesic

cooling itch relief

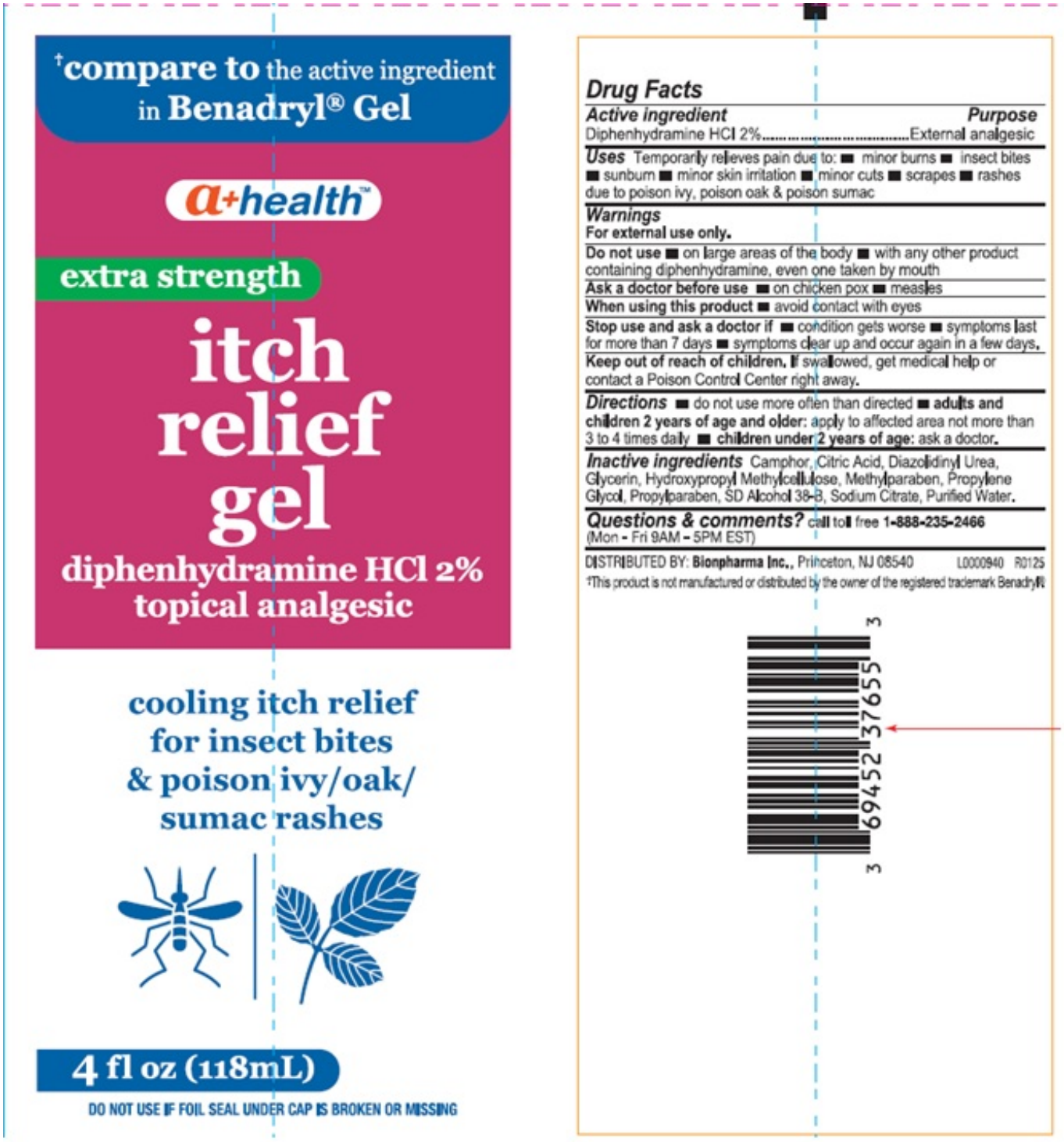
for insect bites

& poison ivy/oak/

sumac rashes

4 fl oz (118 mL)

DO NOT USE IF FOIL UNDER CAP IS BROKEN OR MISSING



A HEALTH ITCH RELIEF GEL

itch relief gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69452-496
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:69452-496-55	118 mL in 1 TUBE; Type 0: Not a Combination Product	06/20/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/20/2025	

Labeler - Bionpharma, Inc. (079637826)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	label(69452-496) , pack(69452-496) , analysis(69452-496) , manufacture(69452-496)

Revised: 6/2025 Bionpharma, Inc.