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

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## A+ Health Itch Relief Gel

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.

For external use only.

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

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.

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

External Analgesic

Diphenhydramine HCl 2%

## Old Labels

†compare to the active ingredient in Benadryl® Gel

a+health™

extra strength

# itch relief gel

**diphenhydramine HCl 2%**
**topical analgesic**

relieves itching & pain associated with insect bites & rashes due to poison ivy, oak & sumac

**4 fl oz (118mL)**

DO NOT USE IF FOIL SEAL UNDER CAP IS BROKEN OR MISSING

3 69452 37655 3

OPEN TO READ COMPLETE PRODUCT INFORMATION

OPEN

## Drug Facts

**Active ingredient** — **Purpose**
Diphenhydramine HCl 2%..... 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, Diazolidinyl Urea, Glycerin, Hydroxypropyl Methylcellulose, Methylparaben, Propylene Glycol, Propylparaben, SD Alcohol 38-B, Sodium Citrate, Purified Water.

**Questions & comments?** call toll free **1-888-235-2466** (Mon - Fri 9AM - 5PM EST)

DISTRIBUTED BY: **Bionpharma Inc.**, Princeton, NJ 08540 L0000589 R03/22
*This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, owner of the registered trademark Benadryl®.

†compare to the active ingredient in Benadryl® Gel

a+health™

extra strength

itch relief gel

diphenhydramine HCl 2%
topical analgesic

cooling relief for insect bites & poison ivy/oak/sumac rashes

4 fl oz (118mL)

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DISTRIBUTED BY: **Bionpharma Inc.**, Princeton, NJ 08540 L0000766 R0624
†This product is not manufactured or distributed by the owner of the registered trademark Benadryl®.

**New Label**


## A HEALTH ITCH RELIEF GEL

itch relief gel gel

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69452-376 |
| **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:69452-376-55 | 118 mL in 1 CONTAINER; Type 0: Not a Combination Product | 11/08/2022 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M017 | 11/08/2022 | |

## Labeler - Bionpharma, Inc. (079637826)

## Registrant - Pharma Nobis, LLC (118564114)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Pharma Nobis, LLC | | 118564114 | manufacture(69452-376) , label(69452-376) , pack(69452-376) , analysis(69452-376) |

Revised: 12/2025 Bionpharma, Inc.