

**WALGREENS STOP ITCH GEL- diphenhydramine hcl gel**  
**Walgreens Co**

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**Walgreens Stop Itch Gel, Diphenhydramine HCl 2%**

Diphenhydramine HCl 2%

Topical Analgesic.

For the temporary relief of pain and itching associated with insect bites, minor burns, sunburn, minor cuts, scrapes, minor skin irritations, and rashes due to poison ivy, oak, and sumac.

**For external use only. Do not use** on large areas of the body, with any other products containing diphenhydramine, even one taken by mouth. **Ask a doctor before use** on chicken pox or measles. **When using this product** avoid contact with eyes. **Stop use and ask a doctor if** condition worsens, if symptoms persist for more than 7 days or clear up and occur again within a few days.

If swallowed, get medical help or contact a Poison Control Center right away.

Do not use more than directed. Adults and children 12 years of age and older, apply to the affected area not more than 3 to 4 times daily. Children under 12 years of age: ask a doctor.

Camphor, Citric Acid, Ethylhexylglycerin, Glycerin, Hydroxypropyl Methylcellulose, Phenoxyethanol, Propylene Glycol, SD Alcohol 38-B, Sodium Citrate, Water.

Walgreens



Compare to the active ingredient in Benedryl®††

Extra Strength

# Stop Itch Gel

DIPHENHYDRAMINE HCl 2%  
TOPICAL ANALGESIC

## FOR SKIN USE ONLY

- Cooling relief for most outdoor itches, insect bites, poison ivy, oak & sumac, mosquito bites, sunburn, minor cuts & scrapes
- Paraben Free
- Fragrance Free

3.5 FL OZ (103 mL)

### Drug Facts

Active ingredient	Purpose
Diphenhydramine HCl 2%.....	Topical analgesic

### Uses

For the temporary relief of 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 products containing diphenhydramine, even one taken by mouth

**Ask a doctor before use** • on chicken pox • on measles

**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 the reach of children.** If product is swallowed, get medical help or contact a Poison Control Center right away.

### Directions

- do not use more than directed
- adults and children 2 years or 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, citric acid, ethylhexylglycerin, glycerin, hydroxypropyl methylcellulose, phenoxyethanol, propylene glycol, SD alcohol 38-B, sodium citrate, water

### Questions or comments?

please call 1-800-925-4733

††This product is not manufactured or distributed by Johnson & Johnson Consumer Inc, owner of the registered trademark Benedryl®

NDC 0000-0000-00

ITEM 000000 W00000-0000-0

DISTRIBUTED BY: WALGREEN CO.

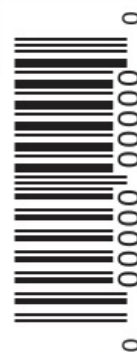
DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

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IMPORTED INGREDIENTS



## WALGREENS STOP ITCH GEL

diphenhydramine hcl gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-1710
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>CAMPHOR (SYNTHETIC)</b> (UNII: 5TJD82A1ET)	
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-1710-03	103 mL in 1 TUBE; Type 0: Not a Combination Product	01/01/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2025	

**Labeler** - Walgreens Co (008965063)

**Registrant** - Derma Care Research Labs, LLC (116817470)

Revised: 9/2024

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