

WALGREENS STOP ITCH GEL- diphenhydramine hcl gel
Walgreens Co

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 Benadryl® Extra Strength Topical Analgesic Itch Stopping Gel^{††}

Itch Relief Gel

DIPHENHYDRAMINE HCl 2% / TOPICAL ANALGESIC

Extra Strength*

- Cooling itch relief associated with sunburn, minor cuts & scrapes, insect bites & poison ivy, oak & sumac
- Paraben free
- Fragrance free

3.5 FL OZ (104 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 chicken pox, poison ivy, sunburn, large areas of the body, broken, blistered, or oozing skin, more often than directed, or with any other product 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

SD alcohol 38-B, camphor, citric acid, ethylhexylglycerin, glycerin, hydroxypropyl methylcellulose, phenoxyethanol, propylene glycol, purified water, sodium citrate

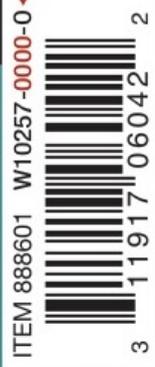
Questions or comments?
please call 1-800-925-4733

*Maximum strength available as an OTC monographed drug.

[†]Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.

^{††}This product is not manufactured or distributed by Kenvue Inc., owner of the registered trademark Benadryl®.

NDC 0363-1710-03 W3ORG0125-F
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DEERFIELD, IL 60015
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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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DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 mL
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Inactive Ingredients

Ingredient Name	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM CITRATE (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: 5/2025

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