WALGREENS STOP ITCH GEL- diphenhydramine hcl gel
Walgreens Co

Walgreens Stop Itch Gel, Diphenhydramine HCl 2%

Diphenhydramine HCl 2%

Topical Analgesic.

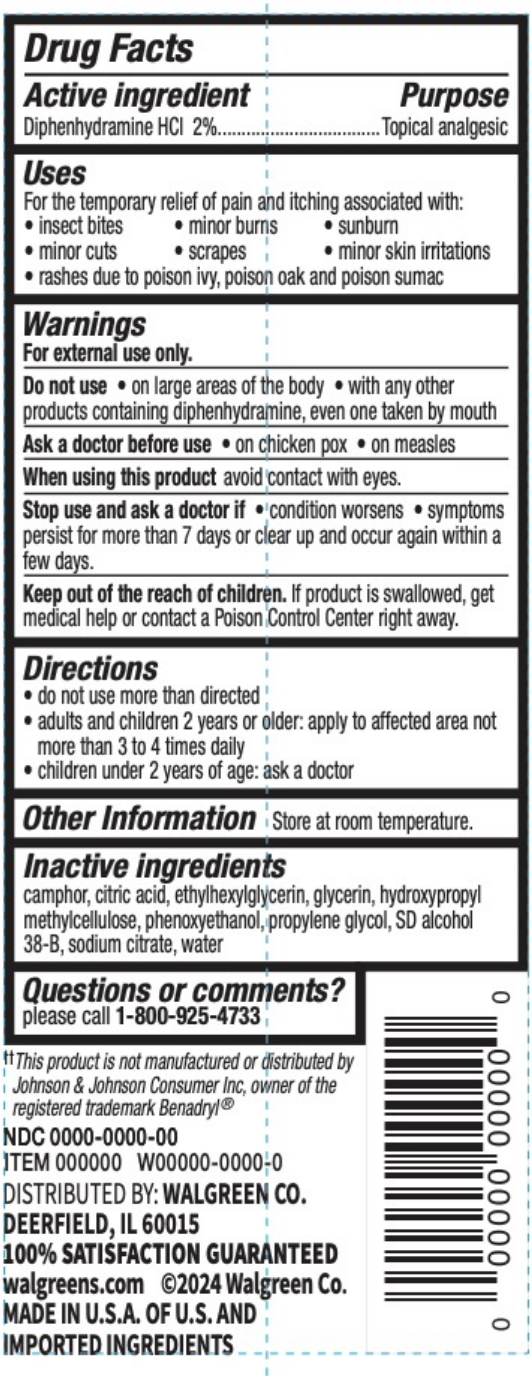
For the temporary relief of pain and itching associated with insect bites, minor burns, sunburn, minor cuts, scrapes, minor skin irritations, and rashes due to poison ivy, oak, and sumac.

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 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**

DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 mL

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: 9/2024 Walgreens Co