

ITCH RELIEF GEL- diphenhydramine hcl gel
Walgreens

Walgreens Itch Relief Gel

Active ingredient

Diphenhydramine HCL 2%

Purpose

Topical Analgesic

Use

Temporarily relieves 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 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 reach of children.

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

Directions

Do not use more than directed.

Adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily.

Other information

Store at room temperature

Inactive ingredients

camphor, diazolidinyl urea, glycerin, hydroxypropyl methylcellulose, methylparaben, propylene glycol, propylparaben, purified water, SD alcohol 40-B, sodium citrate

Questions?

1-800-925-4733

Directions

Apply to affected area not more than 3 to 4 times daily

Tube label

Walgreens

Compare to Benadryl® Extra Strength Topical Analgesic Itch Stopping Gel active ingredient††

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

3.5 FL OZ (103.5 mL)

Drug Facts

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

Use

- temporarily relieves 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 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 reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- do not use more 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

Other Information Store at Room Temperature

Inactive ingredients camphor, diazolidinyl urea, glycerin, hydroxypropyl methylcellulose, methylparaben, propylene glycol, propylparaben, purified water, SD alcohol 40-B, sodium citrate, fragrance.

Questions? 1-800-925-4733

Walgreens Pharmacist Recommended
Walgreens Pharmacist Survey
†† This product is not manufactured or distributed by Johnson & Johnson Corporation, distributor of Benadryl® Extra Strength Topical Analgesic Itch Stopping Gel.

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200 WILMOT RD., DEERFIELD, IL 60015
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ITEM 401632 W10472-1217-F 3 11917 19591 9

ORG1117-F-R1

ITCH RELIEF GEL

diphenhydramine hcl gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0353
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
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
ALCOHOL (UNII: 3K9958V90M)	
HYPROMELLOSE 2208 (15000 MPA.S) (UNII: Z78RG6M2N2)	

Product Characteristics

Color	white (clear)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0353-04	103.5 mL in 1 TUBE; Type 0: Not a Combination Product	05/30/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/30/2018	

Labeler - Walgreens (008965063)**Registrant** - Unipack LLC (116015769)**Establishment**

Name	Address	ID/FEI	Business Operations
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Unipack LLC

009248480

manufacture(0363-0353)

Revised: 2/2024

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