BENADRYL EXTRA STRENGTH ITCH STOPPING- diphenhydramine hydrochloride gel
Johnson & Johnson Consumer Inc.

Benadryl ® EXTRA STRENGTH ITCH STOPPING GEL

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

SD alcohol 38-B, camphor, citric acid, diazolidinyl urea, glycerin, hydroxypropyl methylcellulose, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate

Questions or comments?

call **1-877-717-2824** (toll free) or **215-273-8755** (collect) www.benadryl.com

Distributed by:
JOHNSON & JOHNSON
CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 103 mL Tube Label

Benadryl®

®

Cooling relief for

most outdoor itches

EXTRA STRENGTH

For Skin Use Only

ITCH STOPPING

GEL

Diphenhydramine HCl 2% /

Topical Analgesic

Insect **Poison Ivy,** **Mosquito** **Minor Cuts** **Sunburn**

Bites Oak, Sumac Bites & Scrapes

3.5 FL OZ (103 mL)

BENADRYL EXTRA STRENGTH ITCH STOPPING

diphenhydramine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0242
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0242-3	103 mL in 1 TUBE; Type 0: Not a Combination Product	10/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/01/2013	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 5/2024 Johnson & Johnson Consumer Inc.