BENADRYL ORIGINAL STRENGTH ITCH STOPPING- diphenhydramine hydrochloride and zinc acetate cream
Johnson & Johnson Consumer Inc.

Benadryl ® ORIGINAL STRENGTH ITCH STOPPING CREAM

Drug Facts

Active ingredients	***Purpose***
Diphenhydramine hydrochloride 1%	Topical analgesic
Zinc acetate 0.1%	Skin protectant

Uses

- temporarily relieves pain and itching associated with:
- insect bites
- minor burns
- sunburn
- minor skin irritations
- minor cuts
- scrapes
- rashes due to poison ivy, poison oak, and poison sumac
- dries the oozing and weeping of 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 or does not improve within 7 days
- 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

- protect from excessive heat (40°C/104°F)

Inactive ingredients

cetyl alcohol, diazolidinyl urea, methylparaben, polyethylene glycol monostearate 1000, propylene glycol, propylparaben, purified water

Questions?

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 - 28.3 g Tube Carton

Benadryl ®®

THE HISTAMINE BLOCKER TM

Relief from most outdoor itches

ORIGINAL STRENGTH

For Ages 2+

ITCH STOPPING

CREAM

Diphenhydramine HCl 1%/

Topical Analgesic

Zinc acetate 0.1% /

Skin Protectant

Insect Bites

Poison Ivy,

Oak, Sumac

Mosquito

Bites

Sunburn

Minor Cuts

& Scrapes

NET WT. 1 OZ (28.3g)

BENADRYL ORIGINAL STRENGTH ITCH STOPPING

diphenhydramine hydrochloride and zinc acetate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0625
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	10 mg in 1 g
		1 mg

ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0625-1	1 in 1 CARTON	01/31/2020	
1		28.3 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/31/2020	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2024 Johnson & Johnson Consumer Inc.