GOOD NEIGHBOR PHARMACY ITCH RELIEF EXTRA STRENGTH- diphenhydramine, zinc acetate aerosol, spray
AmerisourceBergen Drug Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Good Neighbor Pharmacy Itch Relief

Drug Facts

Active ingredients

- Diphenhydramine HCI 2.0%
- Zinc Acetate 0.1%

Purpose

- Topical analgesic
- Skin protectant

Uses

Temporarily relieves pain and itching associated with: • insect bites • minor

burns • sunburn • minor cuts • scrapes • minor skin irritations • rashes due to poison

ivy, oak and sumac • dries the oozing and weeping of poison ivy, oak and sumac

Warnings

For external use only.

Flammable:

Do not use while smoking or near heat or flame

Do not use

on large areas of the body • with any other product containing diphenhydramine, even one taken by mouth

When using this product

• keep out of eyes • use only as directed.

• do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120°F.

Ask a doctor before use

• on chicken pox • on measles

Stop use and ask doctor if

• condition worsens • symptoms last more than 7 days

• symptoms clear up and occur again in a few days

Keep out of reach of children.

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

Directions

• shake well • 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 • to apply to face, spray into palm of hand and gently apply

Inactive ingredients

Alcohol Denat., Glycerin, PVP, Tromethamine, Water.

*This product is not manufactured or distributed

by McNeil Consumer Healthcare, owner and

distributor of the Benadryl® trademark.

Distributed By AmerisourceBergen

1300 Morris Drive, Chesterbrook, PA 19087

Visit us at www.goodneighborpharmacy.com

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:46122-114
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 g
Zinc Acetate (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	0.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PVP/VA COPOLYMER (UNII: D9C330MD8B)	
TROMETHAMINE (UNII: 023C2WHX2V)	

WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-114-21	85 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	10/28/2011	

Labeler - AmerisourceBergen Drug Corporation (007914906)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg, LLC		927768135	manufacture(46122-114) , label(46122-114)

Revised: 9/2015 AmerisourceBergen Drug Corporation