SELECT BRAND ITCH RELIEF- diphenhydramine hydrochloride and zinc acetate spray
Select Brand

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.

Select Brand Itch Relief Spray

Drug Facts

Active ingredients	***Purpose***
Diphenhydramine hydrochloride 2%	Topical analgesic
Zinc acetate 0.1%	Skin protectant

Uses

- temporarily relieves pain and itching due to:
  - 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 • oak • 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 do not get into eyes

Stop use and ask a doctor if

- condition worsens
- symptoms last 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: spray on affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Other Information

- store at 20° C to 25° C (68° F to 77° F)

Inactive ingredients

glycerin, povidone, purified water, SD alcohol 40-B, trolamine

Distributed by:

SELECT BRAND DISTRIBUTORS

Pine Bluff, AR 71603 USA, AC(870) 535-3635

PRINCIPAL DISPLAY PANEL

FOR SKIN USE ONLY

select brand

Topical Analgesic

Skin Protectant

ITCH RELIEF SPRAY

*Compare to the active ingredients in

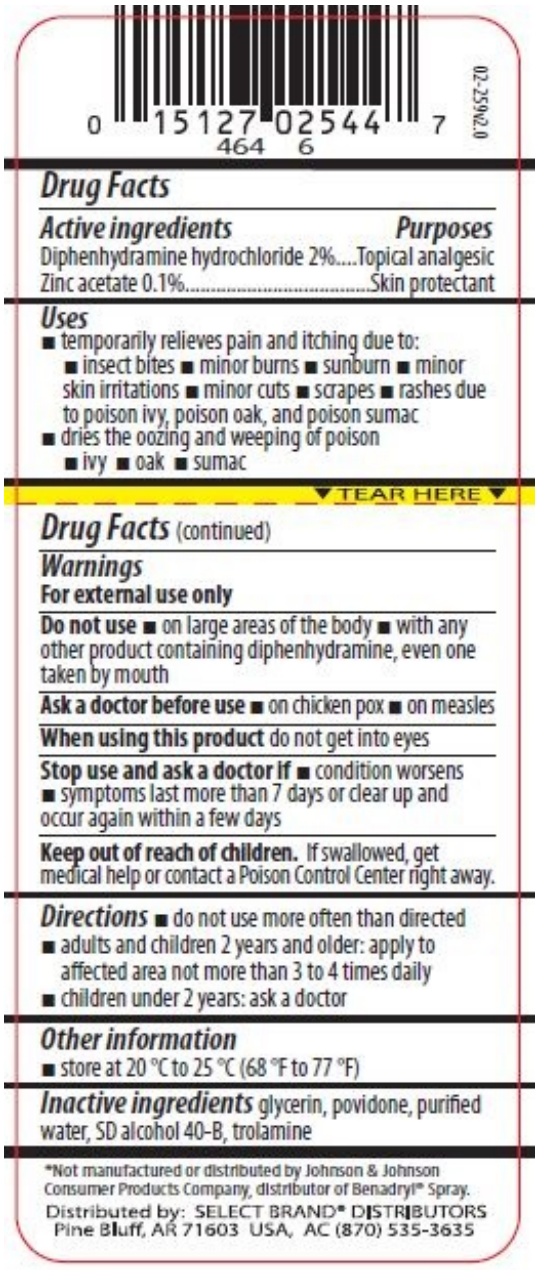
BENADRYL ITCH RELIEF SPRAY

EXTRA STRENGTH

Diphenhydramine HCl 2%

Zinc Acetate 0.1%

2 FL OZ (59 mL)



SELECT BRAND ITCH RELIEF

diphenhydramine hydrochloride and zinc acetate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15127-054
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 10 mL
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	.01 g in 10 mL

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
POVIDONES (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15127-054-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/03/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/11/2012	

Labeler - Select Brand (043562370)

Registrant - Weeks & Leo Co., Inc. (005290028)

Establishment

Name	Address	ID/FEI	Business Operations
Weeks & Leo Co., Inc.		005290028	manufacture(15127-054)

Revised: 1/2015 Select Brand