**CVS ITCH RELIEF- diphenhydramine hcl, zinc acetate spray**
**CVS**

*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.*

---

**CVS Extra Strength Itch Relief Spray**

**Active Ingredient**

Diphenhydramine hydrochloride 2% Topical analgesic

Zinc acetate 0.1% Skin protectant

**Uses**

for

temporary 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

**Directions**

do not use more often than directed

adults and children 2 years and older: apply to affected area not more than 3 to 4 times daily

children under 2 years: ask a doctor

**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 doctor Before use**

on chicken pox

on measles

**When using this product**

do not get into eyes

**Stop Use and ask physician**

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.

**Warning**

for external use only

**Purposes**

Itch relief

**Inactive Ingredients**

Purified Water, Glycerin, SD Alcohol, Povidone, Povidonee (K-30), Trolamine

**Prinicpal dispaly panel**


Diphenhydramine hydrochloride 2%

Topical analgesic

Zinc acetate 0.1% Skin protectant

Relieves Itch and Pain associated with insects bites & rashes due to posinon ivy Oak& sumac

## CVS ITCH RELIEF

diphenhydramine hcl, zinc acetate spray

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69842-952 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ZINC ACETATE** (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37) | ZINC ACETATE | 0.1 g in 100 mL |
| **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M) | DIPHENHYDRAMINE HYDROCHLORIDE | 2 g in 100 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **ALCOHOL** (UNII: 3K9958V90M) | |
| **POVIDONE K30** (UNII: U725QWY32X) | |
| **TROLAMINE** (UNII: 9O3K93S3TK) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:69842-952-02 | 59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 04/14/2014 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part348 | 04/14/2014 | |

## Labeler - CVS (062312574)

## Registrant - Weeks & LEO, Inc. (005290028)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Weeks & Leo | | 005290028 | manufacture(69842-952) |

Revised: 4/2019 CVS