

**CVS EXTRA STRENGTH ITCH STOPPING- diphenhydramine hydrochloride and zinc acetate cream**  
**CVS**

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**CVS Extra Strength Itch Stopping Cream Drug Facts**

**Active Ingredients**

Diphenhydramine hydrochloride 2%

Zinc acetate 0.1%

**Purpose**

Diphenhydramine hydrochloride.....Topical anesthetic

Zinc acetate.....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 due to 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 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

**Other Information**

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

**Inactive Ingredients**

cetyl alcohol, diazolidinyl urea, glyceryl stearate, methylparaben, PEG-40 stearate, PEG-100 stearate, propylene glycol, propylparaben, purified water

**Package Label**



relieves pain from insects bites & skin irritation

**Extra Strength  
ITCH STOPPING CREAM**

Topical Analgesic / Skin Protectant

Net wt. 1 oz (28 g)

**CVS EXTRA STRENGTH ITCH STOPPING**

diphenhydramine hydrochloride and zinc acetate cream

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69842-321
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	20 mg in 1 g
<b>ZINC ACETATE</b> (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	1 mg in 1 g

## Inactive Ingredients

Ingredient Name	Strength
<b>DIAZOLIDINYL UREA</b> (UNII: H5RIZ3MPW4)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PEG-40 STEARATE</b> (UNII: ECU18C66Q7)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-321-28	1 in 1 CARTON	03/17/2014	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69842-321-56	1 in 1 CARTON	03/17/2014	
2		56 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	03/17/2014	

**Labeler** - CVS (062312574)

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

## Establishment

Name	Address	ID/FEI	Business Operations
Weeks & Leo Co., Inc.		005290028	manufacture(69842-321)

