CVS REGULAR STRENGTH ITCH STOPPING- diphenhydramine hydrochloride, zinc acetate cream
CVS

CVS Regular Strength Itch Stopping Cream Drug Facts

Active Ingredients

Diphenhydramine hydrochloride 2%

Zinc acetate 0.1%

Purpose

Diphenhydramine hydrochloride.....Topical anagesic

Zinc acetate.....Skin protectant

Uses

temporarliy 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 20º C to 25º C (68º F to 77º 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

Topical Analgesic

Skin Protectant

ITCH Itch relief CREAM

compare to the active ingredients in Benadryl ® Orignial strength itch stopping cream

Relieves Pain & Itch fast

Itch stopping Cream

CVS REGULAR STRENGTH ITCH STOPPING

diphenhydramine hydrochloride, zinc acetate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-621

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

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-621-01	1 in 1 CARTON	03/17/2014	
1		28 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(69842-621)