UP AND UP ANTI ITCH- hydrocortisone cream Target 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.

Target Corporation Anti-Itch Cream Drug Facts

Active ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Uses

- temporarily relieves itching associated with minor skin irritations, inflammation, and rashes due to:
- eczema
- psoriasis
- poison ivy, oak, sumac
- insect bites
- detergents
- jewelry
- cosmetics
- soaps
- seborrheic dermatitis
- temporarily relieves external anal and genital itching
- other uses of this product should be only under the advice and supervision of a doctor

Warnings

For external use only

Do not use

- in the genital area if you have a vaginal discharge. Ask a doctor.
- for the treatment of diaper rash. Ask a doctor.

When using this product

- avoid contact with the eyes
- do not use more than directed unless told to do so by a doctor
- do not put directly into the rectum by using fingers or any mechanical device or applicator

Stop use and ask a doctor if

condition worsens

- symptoms persist for more than 7 days or clear up and occur again within a few days, and do not begin use of any other hydrocortisone product unless you have asked a doctor
- rectal bleeding occurs

Keep out of reach of children.

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

Directions

- for itching of skin irritation, inflammation, and rashes:
- 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: do not use, ask a doctor
- for external anal and genital itching, adults:
- when practical, clean the affected area with mild soap and warm water and rinse thoroughly
- gently dry by patting or blotting with toilet tissue or a soft cloth before applying
- apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: ask a doctor

Other information

• store at 20°-25°C (68°-77°F)

Inactive ingredients

aloe barbadensis leaf juice, aluminum sulfate, calcium acetate, cetearyl alcohol, cetyl alcohol, cholecalciferol, dextrin, glycerin, isopropyl palmitate, light mineral oil, maltodextrin, methylparaben, propylene glycol, propylparaben, purified water, retinyl palmitate, sodium cetearyl sulfate, sodium lauryl sulfate, tocopherol, white petrolatum, white wax, zea mays (corn) oil

Questions?

Call 1-800-910-6874

Principal Display Panel

Compare to active ingredient in Cortizone - 10® Plus

anti-itch cream

maximum strength

hydrocortisone 1%

NET WT 1 OZ (28 g)

anti-itch cream

maximum strength

hydrocortisone 1%

relieves itch fast

plus 10 moisturizers





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 - insect bites detergents jewelry cosmetics soaps seborrheic dermatitis
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Drug Facts (continued)

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Questions? Call 1-800-910-6874



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UP AND UP ANTI ITCH

hydrocortisone cream

Product Information					
Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:11673-973		
Route of Administration	TOPICAL	DEA Schedule			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
HYDRO CORTISONE (HYDRO CORTISONE)	HYDROCORTISONE	1 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF	
ALUMINUM SULFATE	
CALCIUM ACETATE	
CETOSTEARYL ALCOHOL	
CETYL ALCOHOL	
CHOLECALCIFEROL	
GLYCERIN	
ISOPROPYL PALMITATE	
LIGHT MINERAL OIL	
MALTO DEXTRIN	
METHYLPARABEN	
PROPYLENE GLYCOL	
PROPYLPARABEN	
WATER	
VITAMIN A PALMITATE	
SODIUM CETOSTEARYL SULFATE	
SODIUM LAURYL SULFATE	
TOCOPHEROL	
PETROLATUM	
WHITE WAX	
CORN OIL	

Product Characteristics			
Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packagin	g			
# Ite	m Code	Package Description	Marketing Start Date	Marketing End Date

1 NDC:11673-973-64	1 in 1 CARTON			
1	28 g in 1 TUBE			
Marketing Information				
Marketing Category	Application Number or Monog	raph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348		03/18/2013	

Labeler - Target Corporation (006961700)

Revised: 9/2013 Target Corporation