

UP AND UP HYDROCORTISONE 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 Hydrocortisone 1% 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 only be 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. Consult a doctor.
- for the treatment of diaper rash. Consult 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 (1-800-222-1222).

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, consult 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: consult a doctor

Other information

- store at 20-25°C (68-77°F)

Inactive ingredients

water, cetearyl alcohol, ceteareth-20, cetyl palmitate, glycerin, isopropyl myristate, isostearyl neopentanoate, methylparaben, aloe barbadensis leaf juice

Questions?

Call 1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Cortizone-10[®]

maximum strength

hydrocortisone 1% anti-itch cream with soothing aloe

Compare to active ingredient in Cortizone-10[®]

maximum strength

hydrocortisone 1% anti-itch cream with soothing aloe

#1 doctor recommended anti-itch active ingredient

relieves itch fast

NET WT 1 OZ (28 g)

NDC 11673-842-64



Compare to active ingredient in Cortizone•10®*

maximum strength
hydrocortisone 1% anti-itch cream
with soothing aloe



Compare to active ingredient in Cortizone•10®*

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hydrocortisone 1% anti-itch cream
with soothing aloe

#1 doctor recommended anti-itch active ingredient†
relieves itch fast

NET WT 1 OZ (28 g)



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*This product is not manufactured or distributed by Chattem, Inc., distributor of Cortizone•10®.

†Of U.S. Physicians surveyed by an independent market research firm.

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Drug Facts (continued)

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UP AND UP HYDROCORTISONE ANTI ITCH

hydrocortisone cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:11673-842

Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	HYDROCORTISONE (UNII: W14X0X7BPJ) (HYDROCORTISONE - UNII:W14X0X7BPJ)	HYDROCORTISONE	1 g in 100 g	
Inactive Ingredients				
	Ingredient Name	Strength		
	ALOE VERA LEAF (UNII: ZY81Z83H0X)			
	POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)			
	CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)			
	CETYL PALMITATE (UNII: 5ZA2S6B08X)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
	METHYLPARABEN (UNII: A218C7H9T)			
	WATER (UNII: 059QF0K00R)			
	ISO STEARYL NEOPENTANOATE (UNII: 411THY156Q)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-842-64	1 in 1 CARTON	02/14/2020	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:11673-842-00	2 in 1 CARTON	02/14/2020	
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 not final	part348	02/14/2020	

Labeler - Target Corporation (006961700)

Revised: 10/2020

Target Corporation