

UP AND UP HYDROCORTISONE ANTI ITCH- hydrocortisone cream
Target Corporation

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, cetareth-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®*

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)



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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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- avoid contact with eyes
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- 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 another hydrocortisone product unless you have asked a doctor
- rectal bleeding occurs

Drug Facts (continued)

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Directions

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Other information

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Inactive ingredients

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

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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: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	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: A2I8C7HI9T)	
WATER (UNII: 059QF0KO0R)	
ISOSTEARYL 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	09/30/2022
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:11673-842-00	2 in 1 CARTON	02/14/2020	06/30/2026
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	02/14/2020	06/30/2026

Labeler - Target Corporation (006961700)

