DG HEALTH HYDROCORTISONE- hydrocortisone cream Dolgencorp, LLC

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.

Dolgencorp, LLC Hydrocortisone Cream 1% Drug Facts

Active ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Uses

- temporarily relieves itching associated with minor skin irritations, inflammation, and rashes due to:
- eczema
- insect bites
- poison ivy, oak, or sumac
- soaps
- detergents
- cosmetics
- jewelry
- seborrheic dermatitis
- psoriasis
- 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 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: ask a doctor
- for external anal and genital itching, adults:
- when practical, cleanse 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

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

Questions or comments?

1-888-309-9030

Principal Display Panel

DG™ |health

Compare to the active ingredient of Cortaid®

Maximum Strength

Hydrocortisone Cream 1%

Anti-Itch Cream

Helps skin appear visibly healthier from:

- Redness & dryness
- Flaking & scaling
- Cracking

#1 Doctor recommended Anti-Itch active ingredient

NET WT. 2 OZ (56 g)





Compare to the active ingredient of Cortaid®*

Maximum Strength

Hydrocortisone Cream 1%

Anti-Itch Cream

Helps skin appear visibly healthier from:

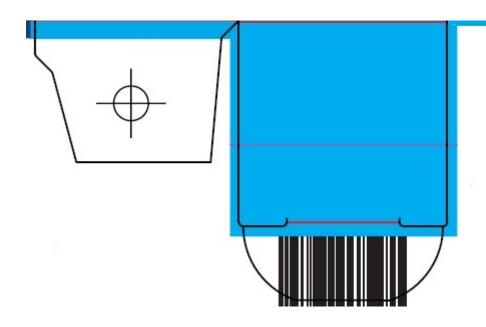
- Redness & dryness
- Flaking & scaling
- Cracking

#1 Doctor recommended Anti-Itch active ingredient

NET WT. 2 0Z (56 g)

Irocortisone Cream 1%

Anti-Itch Cream





DG HEALTH HYDROCORTISONE

hydrocortisone cream

Trydrocordsone cream					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Sou	ırce)	NDC:559	10-541
Route of Administration	TOPICAL				
Active Ingredient/Active Moiety					
Ingre	edient Name		Basis of St	rength	Strength

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)				
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)				
CETYL PALMITATE (UNII: 5ZA2S6B08X)				
GLYCERIN (UNII: PDC6A3C0OX)				
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)				
ISOSTEARYL NEOPENTANOATE (UNII: 411THY156Q)				
METHYLPARABEN (UNII: A218C7H19T)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				

Product Characteristics				
Color	WHITE	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:55910-541- 16	1 in 1 CARTON	01/14/2011				
1		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	01/14/2011		

Labeler - Dolgencorp, LLC (068331990)

Revised: 3/2022 Dolgencorp, LLC