#### EQUALINE ANTI ITCH- hydrocortisone cream United Natural Foods, Inc. dba UNFI

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.

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## SuperValu Inc. 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 or comments?

# 1-877-932-7948

## **Principal Display Panel**

EQUALINE®

compare to Maximum Strength Cortizone•10<sup>®</sup> active ingredient

relieves itch fast

skin irritation – rashes inflammation & redness insect bites eczema & psoriasis plus soothing aloe #1 doctor recommended itch relief active ingredient lasts 10 hours† †when used as directed maximum strength anti-itch cream 1% hydrocortisone anti-itch cream plus soothing aloe NET WT 2 OZ (56g)





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hydrocortisone cream						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:411	DC:41163-319	
Route of Administration	TOPICAL					
Active Ingredient/Active	Moiety					
Ingredient Name Ba			<b>Basis of Stre</b>	asis of Strength Stre		
HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) HYDROCORTISO				NE	1 g in 100 g	
Inactive Ingredients						
Inactive Ingredients	Ingredient Name			S	itrength	
-	•			S	itrength	
Inactive Ingredients ALOE VERA LEAF (UNII: ZY81Z83 POLYOXYL 20 CETOSTEARYL E	H0X)			S	itrength	
ALOE VERA LEAF (UNII: ZY81Z83	HOX) THER (UNII: YRC528SWUY)			S	Strength	
ALOE VERA LEAF (UNII: ZY81Z83 Polyoxyl 20 cetostearyl e	HOX) THER (UNII: YRC528SWUY) 2DMT128M1S)			S	strength	
ALOE VERA LEAF (UNII: ZY81Z83 POLYOXYL 20 CETOSTEARYL E CETOSTEARYL ALCOHOL (UNII: 2	HOX) THER (UNII: YRC528SWUY) 2DMT128M1S)			S	Strength	
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Pa	Packaging						
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:41163-319- 16	1 in 1 CARTON	11/17/2010				
1		56 g in 1 TUBE; Type 0: Not a Combination Product					
2	NDC:41163-319- 64	1 in 1 CARTON	11/18/2010				
2		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			
01 fin	FC monograph not Ial	part348	11/17/2010				

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

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United Natural Foods, Inc. dba UNFI