# HYDROCORTISONE WITH ALOE- hydrocortisone cream Taro Phamaceuticals U.S.A., Inc.

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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#### Hydrocortisone 1% with Aloe

**Drug Facts** 

### **Active ingredient**

Hydrocortisone 1%

# Purpose

Anti-itch cream

#### Uses

- temporary relief of itching associated with minor skin irritations and rashes due to
  - eczema
  - insect bites
  - poison ivy, poison oak, or poison sumac
  - soaps
  - detergents
  - cosmetics
  - jewelry
  - seborrheic dermatitis
  - psoriasis
  - external genital and anal 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 eyes
- by putting this product into the rectum by using fingers or any mechanical device or applicator

# Ask a doctor before use if you have

- a vaginal discharge
- rectal bleeding
- diaper rash

When using this product consult a doctor before exceeding recommended dosage

#### Stop use and ask a doctor if

- condition gets worse
- condition persists for more than 7 days
- condition clears up and occurs again within a few days. Do not begin to use any other

hydrocortisone product unless you have consulted a doctor.

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

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 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 application of this product

Children under 12 years of age: consult a doctor

#### Other information

- To open: unscrew cap, pull tab to remove foil seal, and screw cap back onto tube
- store at room temperature
- see carton or tube crimp for lot number and expiration date

## **Inactive ingredients**

aloe barbadensis, cetearyl alcohol/sodium lauryl sulfate/sodium cetearyl sulfate, citric acid, glycerin, glyceryl stearate, methylparaben, mineral oil, paraffin, propylparaben, purified water, stearyl alcohol

#### Questions?

Call 1-866-923-4914

Distributed by:

Taro Pharmaceuticals U.S.A., Inc.

Hawthorne, NY 10532

#### PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

Itch and Rash Relief

MAXIMUM STRENGTH

Hydrocortisone 1% Cream Antipruritic (Anti-Itch)

With Aloe

NET WT 1 oz (28.4 g)



#### HYDROCORTISONE WITH ALOE

hydrocortisone cream

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Product TypeHUMAN OTC DRUGItem Code (Source)NDC:51672-2013

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Hydrocortisone (UNII: WI4X0X7BPJ) (Hydrocortisone - UNII:WI4X0X7BPJ)	Hydro cortiso ne	1 g in 100 g		

SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)  SODIUM LAURYL SULFATE (UNII: 368GB5141J)  CITRIC ACID MO NO HYDRATE (UNII: 2968PHW8QP)  GLYCERIN (UNII: PDC6A3C0OX)  GLYCERYL MO NO STEARATE (UNII: 230OU9 XXE4)  METHYLPARABEN (UNII: A218C7H19T)  MINERAL O IL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 19O0E3H2ZE)  PRO PYLPARABEN (UNII: Z8IX2SC1OH)  WATER (UNII: 059QF0KO0R)	Inactive Ingredients			
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)  SODIUM LAURYL SULFATE (UNII: 368GB5141J)  CITRIC ACID MO NO HYDRATE (UNII: 2968PHW8QP)  GLYCERIN (UNII: PDC6A3C0OX)  GLYCERYL MO NO STEARATE (UNII: 230OU9 XXE4)  METHYLPARABEN (UNII: A218C7H19T)  MINERAL O IL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 19O0E3H2ZE)  PRO PYLPARABEN (UNII: Z8IX2SC1OH)  WATER (UNII: 059QF0KO0R)	Ingredient Name	Strength		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)  CITRIC ACID MONO HYDRATE (UNII: 2968PHW8QP)  GLYCERIN (UNII: PDC6A3C0OX)  GLYCERYL MONOSTEARATE (UNII: 230OU9 XXE4)  METHYLPARABEN (UNII: A218C7H19T)  MINERAL O IL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 1900E3H2ZE)  PROPYLPARABEN (UNII: Z8IX2SC1OH)  WATER (UNII: 059QF0KO0R)	CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)  GLYCERIN (UNII: PDC6A3C0OX)  GLYCERYL MONOSTEARATE (UNII: 2300U9 XXE4)  METHYLPARABEN (UNII: A218C7H19T)  MINERAL OIL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 1900E3H2ZE)  PROPYLPARABEN (UNII: Z8IX2SC1OH)  WATER (UNII: 059QF0KO0R)	SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)			
GLYCERIN (UNII: PDC6A3C0OX) GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)  METHYLPARABEN (UNII: A2I8C7HI9T)  MINERAL O IL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 19O0E3H2ZE)  PROPYLPARABEN (UNII: Z8IX2SC1OH)  WATER (UNII: 059QF0KO0R)	SO DIUM LAURYL SULFATE (UNII: 368GB5141J)			
GLYCERYL MONOSTEARATE (UNII: 230 O U9 XXE4)  METHYLPARABEN (UNII: A2 I8 C7 HI9 T)  MINERAL O IL (UNII: T5L8 T28 FGP)  PARAFFIN (UNII: 19 O 0 E 3 H2 Z E)  PROPYLPARABEN (UNII: Z8 IX2S C 1 O H)  WATER (UNII: 0 59 Q F 0 K O 0 R)	CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)			
METHYLPARABEN (UNII: A218C7H19T)  MINERAL O IL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 1900E3H2ZE)  PROPYLPARABEN (UNII: Z8IX2SC1OH)  WATER (UNII: 059QF0KO0R)	GLYCERIN (UNII: PDC6A3C0OX)			
MINERAL OIL (UNII: T5L8T28FGP)  PARAFFIN (UNII: 19 O0 E3H2ZE)  PROPYLPARABEN (UNII: Z8 IX2SC1OH)  WATER (UNII: 059QF0 KO0R)	GLYCERYL MONOSTEARATE (UNII: 230 OU9 XXE4)			
PARAFFIN (UNII: 1900E3H2ZE) PROPYLPARABEN (UNII: Z8IX2SC1OH) WATER (UNII: 059QF0KO0R)	METHYLPARABEN (UNII: A218 C7H19 T)			
PROPYLPARABEN (UNII: Z8IX2SC1OH) WATER (UNII: 059QF0KO0R)	MINERAL OIL (UNII: T5L8T28FGP)			
WATER (UNII: 059QF0KO0R)	PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E )			
	PROPYLPARABEN (UNII: Z8IX2SC1OH)			
	WATER (UNII: 059QF0KO0R)			
STEARYL ALCOHOL (UNII: 2KR8914H1Y)	STEARYL ALCOHOL (UNII: 2KR89I4H1Y)			

Packaging					
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
1	NDC:51672-2013-1	1 in 1 CARTON	08/23/1995		
1		14.2 g in 1 TUBE; Type 0: Not a Combination Product			
2	NDC:51672-2013-2	1 in 1 CARTON	08/23/1995		
2		28.4 g in 1 TUBE; Type 0: Not a Combination Product			
3	NDC:51672-2013-3	1 in 1 CARTON	07/17/2020		
3		60 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	08/23/1995		

# **Labeler** - Taro Phamaceuticals U.S.A., Inc. (145186370)

Establishment					
Name	Address	ID/FEI	Business Operations		
Taro Phamaceuticals Inc.		206263295	MANUFACTURE(51672-2013)		