SUNMARK HYDROCORTISONE WITH ALOE MAXIMUM STRENGTHhydrocortisone cream Strategic Sourcing Services LLC

Sunmark ™Hydrocortisone with Aloe Maximum Strength

Drug Facts

Active ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Uses

for the temporary relief of itching associated with minor skin irritations, inflammation and rashes due to:

- eczema
- seborrheic dermatitis
- psoriasis
- insect bites
- poison ivy, oak, sumac
- soaps
- detergents
- cosmetics
- jewelry
- 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
- avoid contact with the eyes

Stop using this product and ask a doctor

- if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days
- before you begin using any other hydrocortisone product

Do not use this product and ask a doctor

- if you have a vaginal discharge
- before treating diaper rash

before using on children under 2 years of age

For External Anal Itching Users

- do not exceed the recommended daily dosage unless directed by a doctor
- in case of bleeding, consult a doctor promptly
- do not put this product into the rectum by using fingers or any mechanical device or applicator
- children under 12 years of age: consult a doctor

Keep this and all drugs out of the reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Before using any medication, read all label directions. Keep this carton. It contains important information.

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, ask a doctor

For External Anal Itching Users

- 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

- unscrew cap, pull tab to remove foil seal, and screw cap back onto tube
- if seal has been broken, do not use this product. Return product to the store where you bought it.
- store at controlled room temperature 59°-86°F
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

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

Distributed by McKesson One Post Street San Francisco, CA 94104

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

sunmark ™

hydrocortisone cream 1%

Antipruritic (Anti-Itch)
MAXIMUM STRENGTH WITH ALOE

NET WT 1 OZ (28.4 g)



NDC 49348-521-72

8

hydrocortisone cream 1%



MAXIMUM STRENGTH WITH ALOE



- Eczema Seborrheic Dermatitis Psoriasis
- Insect Bites Poison Ivy Poison Oak Poison Sumac
 - · External Genital and Anal Itching
 - Soaps Cosmetics Detergents Jewelry

<mark>sun</mark>mark_™

hydrocortisone cream 1%

Antipruritic (Anti-Itch)

sun mark_™

MAXIMUM STRENGTH WITH ALOE

NET WT 1 OZ (28.4 a)

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Drug Facts (continued)



Purpose hoti-itch

Another Quality Product
Distributed by MicKesson
One Post Street
San Francisco, CA 94104
Money Back Guarantee
Please visit us at
Please visit us at

WCKE220N

manufactured or distributed by Pfizer, Inc. or its

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- For Exhemal Anal Itching Users:

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Active ingredient Hydrocortisone 1%.....

Drug Facts

(Astl-itnA) sitirurqitnA

hydrocortisone cream 1%

LPK-4639 1003-0 M124

SUNMARK HYDROCORTISONE WITH ALOE MAXIMUM STRENGTH

Product Information

HUMAN OTC DRUG Item Code (Source) NDC:49348-521 **Product Type**

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)		
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
GLYCERIN (UNII: PDC6A3C0OX)		
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
MINERAL OIL (UNII: T5L8T28FGP)		
PARAFFIN (UNII: 1900E3H2ZE)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
WATER (UNII: 059QF0KO0R)		
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)		

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:49348-521- 72	1 in 1 CARTON	05/01/2014	07/31/2026			
1		28.4 g in 1 TUBE; Type 0: Not a Combination Product					
2	NDC:49348-521- 78	1 in 1 CARTON	05/21/2014	07/31/2026			
2		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 Drug	M017	08/23/1995	07/31/2026		

Registrant - Taro Phamaceuticals U.S.A., Inc. (145186370)

Establishment						
Name	Address	ID/FEI	Business Operations			
Taro Phamaceuticals Inc.		206263295	manufacture(49348-521)			

Revised: 10/2024 Strategic Sourcing Services LLC