

HYDROCORTISONE WITH ALOE MAXIMUM STRENGTH- hydrocortisone cream
NuCare Pharmaceuticals, 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.

HYDROCORTISONE WITH ALOE MAXIMUM STRENGTH

Drug Facts

Active ingredient

Hydrocortisone, USP 1%

Purpose

Anti-itch

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

- eczema •insect bites •poison ivy •poison oak •poison sumac •soaps
- jewelry •detergents •cosmetics •psoriasis •seborrheic dermatitis
- for external genital, feminine 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

- for external feminine itching 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 begin the use of any other hydrocortisone product unless directed by a doctor
- for external anal itching:
- do not use more than directed unless directed by a doctor
- do not put this product into the rectum by using fingers or any mechanical device or applicator

Stop use and ask doctor if

- symptoms last for more than 7 days •the condition gets worse •symptoms clear up and occur again in a few days
- rectal bleeding occurs, consult doctor promptly

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, clean the affected area with mild soap and warm water, 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

- Store at room temperature 59°-86°F (15°-30°C). Protect from freezing.
- Before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredients

aloe vera concentrate, cetyl alcohol, glyceryl stearate, isopropyl myristate, methylparaben, polyoxyl 40 stearate, polysorbate 60, propylene glycol, propylparaben, purified water, sorbic acid, sorbitan monostearate, stearyl alcohol, white wax. May contain citric acid or sodium citrate solution to adjust pH.

Questions?

1-800-432-8534 between 9 am and 4 pm EST, Monday – Friday.

PRINCIPAL DISPLAY PANEL

The image shows the principal display panel for Hydrocortisone 1% Cream. At the top, the NuCare Pharmaceuticals, Inc. logo and name are displayed. The product name "Hydrocortisone 1%" is prominently featured in a large font, with "1oz Cream" below it. The NDC number 68071-3197-1 is shown above the product name. To the right, there are two identical blocks of product information, each containing the product name, lot number (000000), NDC number (68071-3197-01), and MFR NDC number (0472-0339-56) with an expiration date of 00-00. A QR code is located below the second block of information, with GTIN 00368071319712, Serial# 0000000002, Exp. Date 00-00, and LOT# 000000. A warning to call a doctor for medical advice about side effects is also present. On the left side, there is a barcode and the text "Apply every _____ hours _____ times a day." Below the barcode is the number 880713197011*000000*000000. The bottom of the panel features a warning: "WARNING: KEEP OUT OF REACH OF CHILDREN" and "STORE AT CONTROLLED TEMPERATURE 59-86°F." The product number R0279001 is also visible.

HYDROCORTISONE WITH ALOE MAXIMUM STRENGTH

hydrocortisone cream

Product Information

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:68071-3197(NDC:0472-0339) |
|--------------|----------------|--------------------|-------------------------------|
|--------------|----------------|--------------------|-------------------------------|

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|--------------|
| HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ) | HYDROCORTISONE | 1 g in 100 g |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| CETYL ALCOHOL (UNII: 936JST6JCN) | |
| GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4) | |
| ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS) | |
| METHYLPARABEN (UNII: A2I8C7HI9T) | |
| POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I) | |
| POLYSORBATE 60 (UNII: CAL22UVI4M) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| PROPYLPARABEN (UNII: Z8IX2SC1OH) | |
| WATER (UNII: 059QF0KO0R) | |
| SORBIC ACID (UNII: X045WJ989B) | |
| SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X) | |
| STEARYL ALCOHOL (UNII: 2KR89I4HIY) | |
| WHITE WAX (UNII: 7G1J5DA97F) | |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) | |
| SODIUM CITRATE (UNII: 1Q73Q2JULR) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:68071-3197-1 | 1 g in 1 TUBE; Type 0: Not a Combination Product | 06/14/2017 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part348 | 07/08/1998 | |

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

| Name | Address | ID/FEI | Business Operations |
|------------------------------|---------|-----------|---------------------|
| NuCare Pharmaceuticals, Inc. | | 010632300 | relabel(68071-3197) |