

HYDROCORTISONE- hydrocortisone cream
NuCare Pharmaceuticals, Inc.

Perrigo 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

- store at 20-25°C (68-77°F)

Inactive ingredients

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

Questions or comments?

1-800-719-9260

Principal Display Panel

The image shows the principal display panel for Hydrocortisone 1% 1oz Cream. The panel is white with black text and features a large, stylized 'N' logo in the center. The text includes the manufacturer's name, NuCare Pharmaceuticals, Inc., and the product name, Hydrocortisone 1% 1oz Cream. It also provides the NDC number (68071-5205-1), the product number (R0279001), and the warning: 'WARNING: KEEP OUT OF REACH OF CHILDREN'. The panel includes a barcode and a QR code. The text is arranged in a structured layout with various sections for product information, patient instructions, and contact details.

NuCare Pharmaceuticals, Inc.

NDC: 68071-5205-1
Hydrocortisone 1%
1oz Cream

See manufacturer's label
for full list of ingredients.

Product #: R0279001

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

Hydrocortisone 1%
Lot: 00000 NDC: 68071-5205-01
MFR NDC: 45802-438-03 Exp.: 00-00
Serial# 0000000002

Hydrocortisone 1%
Lot: 00000 NDC: 68071-5205-01
MFR NDC: 45802-438-03 Exp.: 00-00
Serial# 0000000002

GTIN 00368071520514
Serial# 0000000002
Exp. Date 00-00
LOT#: 00000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Rev 01/01/19

Applied every _____ times a day. _____ hours

6807152050110000000000

Manufactured for: 3 6807152051
Padagis Minneapolis, MN 55427

Packaged By:
NuCare Pharmaceuticals, Inc.
Orange, CA 92667

Patient Instructions:

HYDROCORTISONE

hydrocortisone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-5205(NDC:45802-438)
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
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CETYL PALMITATE (UNII: 5ZA2S6B08X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ISOSTEARYL NEOPENTANOATE (UNII: 411THY156Q)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-5205-1	28 g in 1 TUBE; Type 0: Not a Combination Product	03/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M014	04/13/2011	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-5205)

Revised: 7/2025

NuCare Pharmaceuticals, Inc.