

**HYDROCORTISONE- hydrocortisone cream**  
**NuCare Pharmaceuticals, Inc.**

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**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**

# NuCare Pharmaceuticals, Inc.

NDC: 68071-5205-1

## Hydrocortisone 1% 1oz Cream

See manufacturer's label  
for full list of ingredients.

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.

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

Distributed by:  
Padagis Allegan, MI 49010



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Rev 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

Product #: R0279001

STORE AT CONTROLLED TEMPERATURE 68-77°F.

68071520501-1-00000-00000

Apply every \_\_\_\_\_ hours  
\_\_\_\_\_ times a day.

Patent 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
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CETYL PALMITATE (UNII: 5ZA2S6B08X)	
GLYCERIN (UNII: PDC6A3C0OX)	
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: 1/2025

NuCare Pharmaceuticals, Inc.