

**FREDS HYDROCORTISONE WITH ALOE- hydrocortisone cream
FRED'S, INC.**

**Fred's Hydrocortisone Cream 1%
with Aloe**

Drug Facts

Active ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Uses

- temporarily relieves itching associated with minor skin irritations, inflammation, and rashes due to:
 - eczema
 - psoriasis
 - poison ivy, oak, sumac
 - insect bites
 - detergents
 - jewelry
 - cosmetics
 - soaps
 - seborrheic dermatitis
- temporarily relieves external anal and genital 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 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.

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, clean 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

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 powder, cetearyl alcohol/sodium lauryl sulfate/sodium cetearyl sulfate, citric acid, glycerin, glyceryl monostearate, methylparaben, mineral oil, paraffin wax, propylparaben, purified water, sodium lauryl sulfate, stearyl alcohol

Questions?

Call **1-866-923-4914**

**DISTRIBUTED BY: fred's, Inc.
4300 NEW GETWELL RD,
MEMPHIS, TN 38118**

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

fred's
Pharmacy

NDC 55315-181-56

Compare to the

Active Ingredient in
Cortizone•10[®]*

MAXIMUM STRENGTH

Hydrocortisone
Cream 1%

Antipruritic (Anti-Itch)
Itch and Rash Relief

- Eczema & Psoriasis
- Seborrheic Dermatitis
- Insect Bites & Poison Ivy
- External Genital & Anal Itching
- Skin Irritation & Rashes

With Healing Aloe

NET WT 1 oz (28.4g)

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4300 NEW GETWELL RD,
MEMPHIS, TN 38118
www.fredsinc.com

Made in Canada

100% satisfaction
guaranteed

Questions or comments:
1-855-331-FRED (3733)

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NO COPY ON THIS FLAP
FOR LOT # AND EXPIRY
DATE PRINT ONLY

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Hydrocortisone 1%
Anti-Itch

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Questions? Call 1-866-923-4914
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Antipruritic (Anti-Itch)
MAXIMUM STRENGTH

Hydrocortisone
Cream 1%
With Aloe

FREDS HYDROCORTISONE WITH ALOE

hydrocortisone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55315-181
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)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PARAFFIN (UNII: I9O0E3H2ZE)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55315-181-56	1 in 1 CARTON	11/15/2018	
1		28.4 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	11/15/2018	

Labeler - FRED'S, INC. (005866116)**Establishment**

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(55315-181)