

**ITCH AND RASH CREAM- hydrocortisone cream**  
**Wildman Business Group**

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**1% Hydrocortisone Itch Cream**

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, detergents, cosmetics, jewelry, seborrheic dermatitis, psoriasis and

scrapes

KEEP OUT OF REACH OF CHILDREN. IF SWALLOWED, get medical help or

contact a Poison Control Center right

away.

Warnings :

For external use only Do not use for the treatment of diaper rash

Consult a doctor: before use if you have a vaginal discharge (for external feminine itching):

for external itching, do not exceed the recommended daily dosage or if bleeding occurs if condition worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days

When using this product: avoid contact with eyes, do not put this product into rectum by using fingers or any mechanical

Do not use: with any other Hydrocortison product unless you have consulted a doctor

**DIRECTIONS: FOR ADULTS AND CHILDREN 2 YEARS OF AGE AND OLDER-APPLY TO AFFECTED AREA NOT MORE THAN 3 OR 4 TIMES DAILY**

**CHILDREN UNDER 2 YEARS, DO NOT USE, CONSULT A DOCTOR. ADULTS FOR EXTERNAL ANAL ITCHING WHEN PRACTICAL-CLEANSE THE AFFECTED AREA WITH A MILD SOAP AND WARM WATER AND RINCE THOROUGHLY OR BY PATTING AND BLOTTING WITH AN APPROPRIATE CLEANSING PAD. GENTLY DRY BY PATTING OR BLOTTING WITH A SOFT CLOTH BEFORE APPLICATION OF THIS PRODUCT. CHILDREN UNDER 12 YEARS OF AGE-FOR EXTERNAL ANAL ITCHING, CONSULT A DOCTOR**

**INACTIVE INGREDIENTS: CITRIC acid, glycerin, glycerol stearate, methyl paraben, petrolatum, polysorbate 80, propyl**

**paraben, propylene glycol, purified water, sodium citrate, and titanium dioxide.**

**ACTIVE INGREDIENT (IN EACH GRAM)-HYDROCORTISONE 10 MG**

**ANTIPRURITIC (ANTI-ITCH)**

# 1% Hydrocortisone Itch Cream

## 1% Hydrocortisone/Itch Cream

The Provision  
First Aid  
Line™

BY WILDMAN

0.9 g  
(1/32 oz.)

### Drug Facts

<b>Active ingredients</b>	<b>Purpose</b>
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Hydrocortisone 1.0% . . . . Anti-itch

**Uses** For Temporary relief of itching associated with minor skin irritations, inflammation, or rashes. Other uses of product should be only under the advice and supervision of a doctor.

### Warnings

**For external use only**

**Do not use** ■ in eyes ■ for treatment of diaper rash ■ for feminine itching ▶

Manufactured for: Provision by  
Wildman, Warsaw, IN 46580  
Provisionfirstaid.com 866-369-1552

### Drug Facts (continued)

**Stop use, ask a doctor if**  
■ condition worsens or lasts more than 7 days, or clears up and occurs again within a few days. ■ you begin use of any other hydrocortisone product.

**Keep out of reach of children.**  
If swallowed, get medical help or contact a Poison Control Center directly.

**Directions** ■ Apply to affected area no more than 3 to 4 times daily. ■ For children under 2: ask a doctor.

**Inactive ingredients**  
emulsifying wax, ethanol, methylparaben, mineral oil, paraffin, petrolatum, propylparaben, purified water, white wax

## ITCH AND RASH CREAM

hydrocortisone cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84269-4506
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	10 mg in 1 g

## Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ANHYDROUS TRISODIUM CITRATE</b> (UNII: RS7A450LGA)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84269-4506-1	25 in 1 CARTON	09/01/2024	
1		0.9 g in 1 PACKET; 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	09/01/2024	

**Labeler** - Wildman Business Group (016677338)

**Registrant** - Safetec of America Inc. (874965262)

Revised: 9/2024

Wildman Business Group