

QUALITY CHOICE ANTI-ITCH- hydrocortisone cream
United Exchange Corp.Chain Drug Marketing Association 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.

QC Anti-itch Cream with Aloe 1oz 99259 ZDP (2019)

Active ingredient Purpose

Hydrocortisone 1%.....Anti-itch

Uses

- temporarily relieves itching associated with minor skin irritations, inflammation, and rashes due to:
- eczema
- psoriasis
- jewelry
- insect bites
- soaps
- cosmetics
- detergents
- seborrheic dermatitis
- poison ivy, oak, sumac
- temporarily relieves external anal and genital itching
- other uses of this product should only be under the advise 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.
- more than directed unless directed by a doctor

When using this product

- avoid contact with eyes
- do not put directly into the rectum by using fingers or any mechanical device or applicator

Stop use and ask a doctor

- 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

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) 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 a 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

- store at room temperature 20-25°C (68-77°F)

Inactive ingredients

aloe vera leaf, aluminum sulfate octadecahydrate, calcium acetate, monohydrate, cetostearyl alcohol, dextrin, light mineral oil, maltodextrin, methylparaben, petrolatum, propylene glycol, propylparaben, purified water, sodium cetearyl sulfate, sodium lauryl sulfate, white wax

Distributed by: C.D.M.A., Inc.

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Made in China



NDC 63868-693-28

*Compare to the Active Ingredient in Cortizone-10® Anti-Itch Creme

Anti-Itch Cream with Aloe**

Maximum Strength

Hydrocortisone 1% (Anti-Itch)

Temporary Relief of:
Skin Irritation and Rashes | Inflammation and Redness |
Insect Bites and Poison Ivy | Eczema and Psoriasis



Anti-Itch Cream with Aloe**

Maximum Strength

Hydrocortisone 1% (Anti-Itch)



NET WT. 1 oz (28 g)



Anti-Itch Cream®
with Aloe®
Maximum Strength



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Distributed by C.D.M.A., Inc.®
43157 W 9 Mile Rd
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362

MADE IN CHINA

99259-ZDR-BX-R1
0M-MD-ZDP119-18

LOT & EXP.

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 ■ jewelry ■ insect bites ■ soaps ■ cosmetics ■ detergents ■ sebormic dermatitis ■ poison ivy, oak, sumac ■ temporarily relieves external nasal and genital itching

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. ■ more than directed unless directed by a doctor.
When using this product
avoid contact with eyes
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

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 nasal 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
store at room temperature 20-25°C (68-77°F)
Inactive ingredients side wax leaf, aluminum sulfate octadecylhydrate, calcium acetate monohydrate, cetearyl alcohol, stearyl, light mineral oil, methocel k100, myristyl alcohol, propylene glycol, propylparaben, purified water, sodium cetearyl sulfate, sodium lauryl sulfate, white wax

Drug Facts (continued)
If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

OR FOLLOUNDER CAP IS BROKEN OR MISSING, TAMPER EVIDENT DO NOT USE IF CARTON IS OPEN.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

QUALITY CHOICE ANTI-ITCH

hydrocortisone cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:63868-693

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
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
WHITE WAX (UNII: 7G1J5DA97F)	
ICODextrin (UNII: 2NX48Z0A9G)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALUMINIUM SULFATE OCTADECALHYDRATE (UNII: TCS9L00G8F)	
CALCIUM ACETATE MONOHYDRATE (UNII: 7ZA48GIM5H)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-693-28	1 in 1 CARTON	07/10/2019	
1		28 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 not final	part348	07/10/2019	

Labeler - United Exchange Corp.Chain Drug Marketing Association Inc. (011920774)

Revised: 12/2019

United Exchange Corp.Chain Drug Marketing Association Inc.