QUALITY CHOICE HYDROCORTISONE MAXIMUM STRENGTH- hydrocortisone cream
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 Max. Strength Itch Relief Cream 1 oz 99260, 2019

Active ingredient **Purpose**

Hydrocortisone 1%..... Anti-itch

Uses

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

- conditions 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 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 between 20° to 25°C (68°-77°F)
- Lot No. & Exp. Date: see box or see crimp of tube

Inactive ingredients

butylated hydroxytoluene, cetanol, liquid paraffin, methylparaben, polyoxyethylene cetylether, propylene glycol, propylparaben, purified water, sorbitan monostearate, stearyl alcohol

DISTRIBUTED BY C.D.M.A., INC.

43157 W. NINE MILE

NOVI, MI 48376-0995

www.qualitychoice.com

MADE IN KOREA

NDC 63868-596-28

QUALITY CHOICE

*Compare to the active ingredient in CORTAID®

1% Hydrocortisone

Maximum Strength

Temporary Relief of:
Eczema | Poison Ivy, Oak, and Sumac | Insect bites | Psoriasis | Dermatitis

QUALITY CHOICE

1% Hydrocortisone

Maximum Strength

Temporary Relief of:
Eczema | Poison Ivy, Oak, and Sumac | Insect bites | Psoriasis | Dermatitis

1 oz NET WT (28g)

*This product is not manufactured or distributed by Valeant Consumer Pharmaceuticals North America, owner of the registered trademark Cortaid®.

Made in Korea

SATISFACTION 100% QC GUARANTEED

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www.qualitychoice.com
Questions: 248-449-9300

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QUALITY CHOICE

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Maximum Strength

LOT & EXP.

Drug Facts

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QUALITY CHOICE HYDROCORTISONE MAXIMUM STRENGTH

hydrocortisone cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-596
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**
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PARAFFIN (UNII: I9O0E3H2ZE)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-596-28	1 in 1 CARTON	01/16/2015	
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	01/15/2015	

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 12/2019 Chain Drug Marketing Association Inc