QUALITY CHOICE ANTI ITCH- 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 Anti-Itch Cream Intensive Healing 1 oz. 99198 ZDP

Active ingredient Purpose

Hydrocortisone 1%......Anti-itch

Uses

- temporarily relieves itching associated with minor skin irritation, 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 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.
- 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

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 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, .alpha.-tocopherol acetate, benzyl alcohol, cetyl alcohol, dimethicone 350, edetate disodium, gelatin hydrolysate, glycerin, glyceryl stearate/peg-100 stearate, magnesium ascorbyl, phosphate, matricaria chamomilla, flowering top, menthyl lactate,methyl gluceth-20, methylparaben, petrolatum,polysorbate 60, propylene glycol, propylparaben, purified water, steareth-2, steareth-21, stearyl alcohol, vitamin a palmitate

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

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Novi. MI 48375

www.qualitychoice.com

Made in China



QUALITY CHOICE ANTI ITCH hydrocortisone cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:63868-955

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		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
PETROLATUM (UNII: 4T6H12BN9U)			
BENZYL ALCOHOL (UNII: LKG8494WBH)			
CETYL ALCOHOL (UNII: 936JST6JCN)			
GLYCERIN (UNII: PDC6A3C0OX)			
MAGNESIUM ASCORBYL PHOSPHATE (UNII: 0R822556M5)			
STEARETH-21 (UNII: 53J3F32P58)			
POLYSORBATE 60 (UNII: CAL22UVI4M)			
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			
GLYCERYL STEARATE/PEG-100 STEARATE (UNII: RD25J5V947)			
MATRICARIA CHAMOMILLA FLOWERING TOP (UNII: 3VNC7T6Z02)			
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)			
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)			
METHYL GLUCETH-20 (UNII: J3QD0LD11P)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
DIMETHICONE 350 (UNII: 2Y53S6ATLU)			
GELATIN HYDROLYSATE (PORCINE SKIN, MW 3000) (UNII: 0K9R94573C)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
STEARETH-2 (UNII: V56DFE46J5)			

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:63868-955- 28	1 in 1 CARTON	12/10/2019			
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	11/07/2019		

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 9/2022

Chain Drug Marketing Association Inc