CORTIZONE 10 INTENSIVE HEALING FEMININE ITCH RELIEF-
hydrocortisone cream
Chattem, Inc.

Cortizone 10 Intensive Healing Feminine Itch Relief

Maximum Strength Cotizone 10® Feminine Relief Anti-Itch Creme

Drug Facts

Active ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Uses

- temporarily relieves external feminine itching
- other uses of this product should only be under the advice and supervision of a doctor

Warnings

For external use only

When using this product

- avoid contact with eyes
- do not use more than directed unless told to do so by a doctor

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

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- when practical, cleanse the affected area with mild soap and warm water and rinse thoroughly then gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product
- **adults and children 12 years of age and older:** apply a fingertip amount

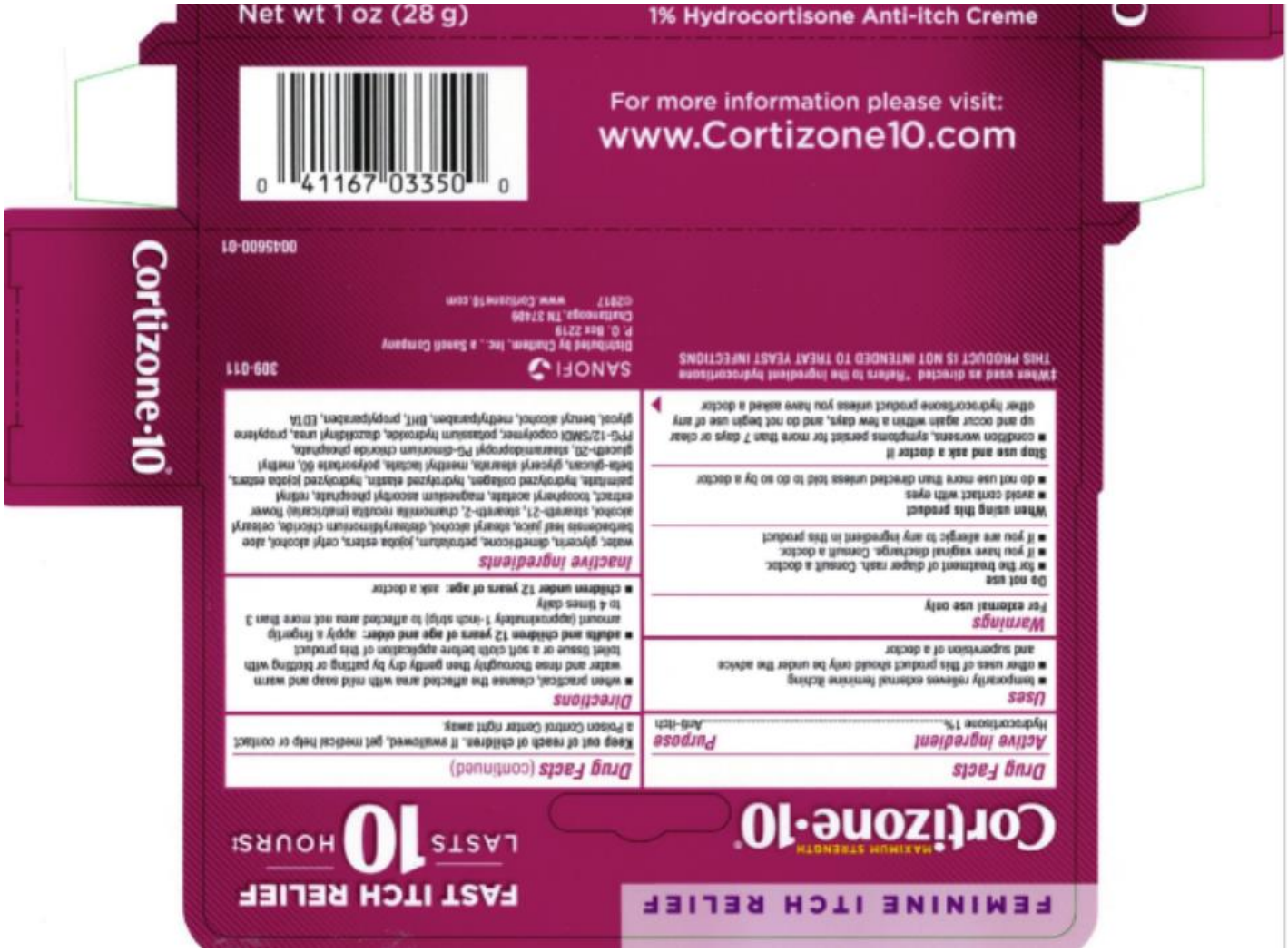
(approximately 1-inch strip) to affected area not more than 3 to 4 times daily
- **children under 12 years of age:** ask a doctor

Inactive ingredients

water, glycerin, dimethicone, petrolatum, jojoba esters, cetyl alcohol, aloe barbadensis leaf juice, stearyl alcohol, distearyldimonium chloride, cetearyl alcohol, steareth-21, steareth-2, chomomilla recutita (matricaria) flower extract, tocopheryl acetate, magnesium ascorbyl phosphate, retinyl palmitate, hydrolyzed collagen, hydrolyzed elastin, hydrolyzed jojoba esters, beta-glucan, glyceryl stearate, menthyl lactate, polysorbate 60, methyl gluceth-20, stearamidopropyl PG-dimonium chloride phosphate, PPG-12/SMDI copolymer, potassium hydroxide, diazolidinyl urea, propylene glycol, benzyl alcohol, methylparaben, BHT, propylparaben, EDTA (309-011)

Principal Display Panel

Maximum Strength
Cotizone 10®
FEMININE ITCH RELIEF
Anti-Itch Crème
For Feminine Itch
1% Hydrocortisone Anti-Itch Creme
Net wt 1 oz (28 g)



CORTIZONE 10 INTENSIVE HEALING FEMININE ITCH RELIEF

hydrocortisone cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41167-0335
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**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PETROLATUM (UNII: 4T6H12BN9U)	
JOJOBA OIL, RANDOMIZED (UNII: 7F0EV20QYL)	

CETYL ALCOHOL (UNII: 936JST6JCN)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARETH-2 (UNII: V56DFE46J5)	
CHAMOMILE (UNII: FGL3685T2X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MAGNESIUM ASCORBYL PHOSPHATE (UNII: 0R822556M5)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
HYDROLYZED BOVINE ELASTIN (BASE; 1000 MW) (UNII: ZR28QKN0WT)	
HYDROLYZED JOJOBA ESTERS (ACID FORM) (UNII: UDR641JW8W)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
STEARAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: W6000VEI5Y)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
EDETIC ACID (UNII: 9G34HU7RV0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0335-0	1 in 1 CARTON	01/01/2014	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:41167-0335-2	1 in 1 CARTON	01/01/2014	01/02/2014
2		56 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:41167-0335-1	1 in 1 CARTON	01/01/2014	01/02/2014
3		14 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	01/01/2014	

Labeler - Chattem, Inc. (003336013)

Revised: 10/2023 Chattem, Inc.