CORTIZONE 10 OVERNIGHT ITCH RELIEF- hydrocortisone cream Chattem, Inc. _____ **Cortizone-10 Overnight Relief** Cortizone-10 Overnight Relief **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 ■ poison ivy, oak, 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

Stop use and ask a doctor if

or applicator

■ 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

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 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

Inactive ingredients

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

Principal Display Panel

OVERNIGHT
ITCH RELIEF
with Lavender Scent
MAXIMUM STRENGTH
Cortizone 10
1% Hydrocortisone Anti-itch Crème
Net wt 1 oz (28 g)



CORTIZONE 10 OVERNIGHT ITCH RELIEF

hydrocortisone cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41167-0108
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
JOJOBA OIL, RANDOMIZED (UNII: 7F0EV20QYL)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CURDLAN (UNII: 6930DL209R)	
HYDROLYSED BOVINE COLLAGEN (ENZYMATIC; 3500 MW) (UNII: 5WE8P977RQ)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y10)	
PETROLATUM (UNII: 4T6H12BN9U)	
HYDROGENATED JOJOBA OIL, RANDOMIZED (UNII: Q47ST02F58)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
MELATONIN (UNII: JL5DK93RCL)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CHAMOMILE (UNII: FGL3685T2X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MAGNESIUM ASCORBYL PHOSPHATE (UNII: 0R822556M5)	
HYDROLYZED BOVINE ELASTIN (BASE; 1000 MW) (UNII: ZR28QKNOWT)	
HYDROLYZED JOJOBA ESTERS (ACID FORM) (UNII: UDR641JW8W)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
STEARAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: W6000VEI5Y)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
EDETIC ACID (UNII: 9G34HU7RV0)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
POTASSIUM HYDROXIDE (UNII: WZ H3C48M4T)	

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:41167-0108-1	1 in 1 CARTON	01/01/2020			
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 Drug	M017	01/01/2020		

Labeler - Chattem, Inc. (003336013)

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