

UEC MEDICAL ANTI-ITCH- hydrocortisone cream
United Exchange Corp.

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.

UEC Medical Hydrocortisone 1% Cream 0.9g NBE Cortaid 20 and 144 ct. - 10393 and 10397 (2018)

Active ingredient Purpose

Hydrocortisone 1%.....Anti-itch

Uses

- temporarily relieves itching associated with minor skin irritations, inflammation, and rashes due to:
- eczema
- insect bites
- poison ivy, oak, or sumac
- soaps
- detergents
- cosmetics
- jewelry
- seborrheic dermatitis
- psoriasis
- temporarily relieves external anal and genital itching
- other uses of this product should be under 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. Ask 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

- 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: 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 tissues 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 20° to 25°C (68° to 77°F)

Inactive ingredients

cetamacrogol 1000, cetostearyl alcohol, chlorocresol, edetate disodium, liquid paraffin, propylene glycol, purified water, sodium metabisulphite, white soft paraffin

Distributed by:

Untied Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 USA



Drug Facts (continued)
device or applicator

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Compare to **CORTAID®** Maximum Strength active ingredient*

Hydrocortisone 1%
ANTI-ITCH CREAM

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

PUSH TO OPEN
Dispense through opening
Do not use if packet is torn or damaged

20 Packets
½ oz (0.9 g) each

Compare to **CORTAID®** Maximum Strength active ingredient*

Hydrocortisone 1%
ANTI-ITCH CREAM

MAXIMUM STRENGTH

HYDROCORTISONE 1%
■ Rapid itch relief from skin irritations and rashes

20 Packets
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Reorder No. 10393
Ointment Made in Israel
Packaged in the U.S.A.



UEC MEDICAL ANTI-ITCH

hydrocortisone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-393
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE (UNII: WI4X0 X7BPJ) (HYDROCORTISONE - UNII:WI4X0 X7BPJ)	HYDROCORTISONE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	

SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETETH-20 (UNII: I835H2IHHX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CHLOROCRESOL (UNII: 36W53O7109)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PARAFFIN (UNII: I9O0E3H2ZE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-393-20	20 in 1 BOX	04/04/2016	
1		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:65923-393-44	144 in 1 BOX	04/04/2016	
2		0.9 g in 1 PACKET; 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	04/04/2016	

Labeler - United Exchange Corp. (840130579)

Revised: 4/2018

United Exchange Corp.