KERA HC- hydrocortisone cream
Clinical Therapeutic Solutions

KERA HC Cream Hydrocortisone 1%

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Active Ingredient

Hydrocortisone-USP 1.00%

Purpose

Anti-Itch

Uses:

Temporarily relieves itching associated with minor skin irritations, inflammations and rashes due to:

- Excessively Dry Skin, Eczema, Insect Bites, Poison Ivy, Oak or Sumac, Soaps, Detergents, Cosmetics, Jewelery, Seborrheic Dermatitis, Psoriasis
- Other uses of this product should only be under the advice and supervisor 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 or symptoms persist for more than 7 days, or if symptoms clear up, and then return

Keep out of reach of children.

If swallowed, contact physician or poison control center immediately.

Store at room temperature (20-25 degrees C) (68-77 degrees F)

Directions:

For itching of skin irritation, inflammation and rashes:

- Apply to affected area not more than 3 to 4 times daily.
- Not for use by children under 2 years of age

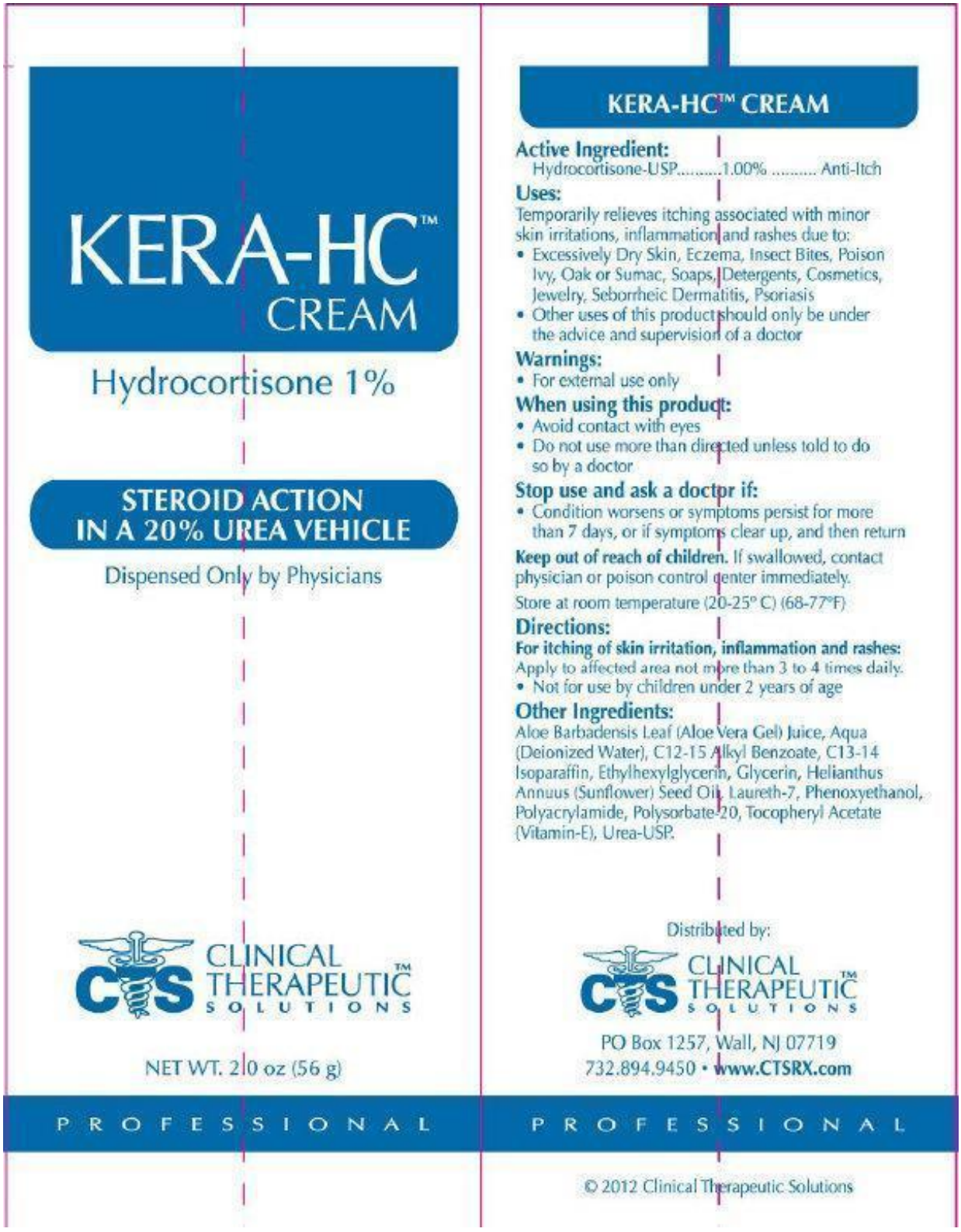
Other Ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), C12-15 Alkyl Benzoate, c13-14 Isoparaffin, Ethylhexylglycerin, Glycerin, Heilianthus Annuus (Sunflower) Seed Oil, Laureth-7, Phenoxyethanol, Polacrylamide, Polysorbate-20, Tocopheryl Acetate (Vitamin E), Urea-USP.

Distributed By:

Clinical Therapeutic Solutions, PO Box 1257, Wall, NJ 07719 www.CTSRX.com

KERA HC Cream Hydrocortisone 1% 2oz/56g (44577-623-02)



KERA HC

hydrocortisone cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44577-623
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
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
LAURETH-7 (UNII: Z95S6G8201)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
UREA (UNII: 8W8T17847W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44577-623-02	56 g in 1 TUBE; Type 0: Not a Combination Product	02/06/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/17/2012	

Labeler - Clinical Therapeutic Solutions (078402750)

Revised: 11/2023 Clinical Therapeutic Solutions