NIZORAL SCALP ITCH RELIEF- hydrocortisone shampoo Kramer Laboratories

Drug Facts

Active ingredient	Purpose
Hydrocortisone	Anti-itch
1.0%	liquid

Uses

- temporarily relieves itching associated with minor skin irritations, inflammation, and rashes due to:
 - eczema
 - psoriasis
 - seborrheic dermatitis
- other uses of this product should be only under the advice and supervision of a doctor

Warnings

For external use only. Use only as directed

Flammable. Keep away from fire, or flame.

Do not use

- for the treatment of diaper rash. Consult a doctor.
- If you are allergic to any ingredient in this product

When using this product

- avoid contact with the 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, do not use this or any other hydrocortisone product unless you have asked a doctor.

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- twist the applicator tip to open.
- if applying after shampooing, towel dry hair before use.

- squeeze bottle gently to avoid excess dripping.
- apply solution directly to scalp and massage in.
- twist applicator tip tightly after use.
- 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: do not use, ask a doctor

Other information

- store product at 20-25°C (68-77°F)
- keep carton for full product information.

Inactive ingredients

alcohol denat., aloe barbadensis leaf juice, dipropylene glycol, disodium EDTA, glycerin, hyaluronic acid, menthol, PEG-12 dimethicone, phenoxyethanol and ethylhexylglycerin, polysorbate-80, tocopheryl acetate, water (aqua)

Questions? 1-800-824-4894

PRINCIPAL DISPLAY PANEL

Nizoral[®]

Scalp Itch Relief

HYDROCORTISONE 1%
ANTI-ITCH LIQUID

RELIEVES SCALP ITCH

from seborrheic dermatitis, psoriasis & eczema

SOOTHES, CALMS & HYDRATES

plus aloe, menthol & hyaluronic acid

MAXIMUM STRENGTH

anti-itch medicine

fragrance-free • greaseless formula

2 fl oz (60mL)

NIZORAL® Scalp Itch Relief

contains maximum strength medicine (Hydrocortisone 1%) to relieve scalp itch. The special medicated solution helps soothe, calm, & hydrate an itchy, irritated scalp; plus it contains aloe, menthol, vitamin E & hyaluronic acid. The tapered bottle tip design helps easily apply the medicated solution that's paraben free, fragrance free, and greaseless.

Specially formulated with:

- √ Aloe
- √ Menthol
- √ Hyaluronic Acid
- √ Vitamin E

Before use, read all label information. If you have a drug reaction, contact a doctor and report it by calling: 1.800.824.4894

KRAMER LABORITORIES

Distributed by: Kramer Laboratories, Inc. Bridgewater, NJ 08807 USA

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NIZORAL SCALP ITCH RELIEF

hydrocortisone shampoo

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55505-222
Route of Administration	TOPICAL			
Active Ingredient/Active	Majahy			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Streng	gth Strength

Hydrocortisone (UNII: W4X0X7BPJ) (Hydrocortisone	- UNII:WI4X0X7BPJ)	Hydrocortisone	10 mg in 1 mL
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Inactive Ingredients		
Ingredient Name	Strength	
Alcohol (UNII: 3K9958V90M)		
Aloe (UNII: V5VD430YW9)		
Dipropylene Glycol (UNII: E107L85C40)		
Edetate Disodium Anhydrous (UNII: 8NLQ36F6MM)		
Glycerin (UNII: PDC6A3C0OX)		
Hyaluronic Acid (UNII: S270N0TRQY)		
Menthol, Unspecified Form (UNII: L7T10EIP3A)		
Peg-12 Dimethicone (300 Cst) (UNII: ZEL54N6W95)		
Phenoxyethanol (UNII: HIE492ZZ3T)		
Ethylhexylglycerin (UNII: 147D247K3P)		
Polysorbate 80 (UNII: 60ZP39ZG8H)		
.AlphaTocopherol Acetate (UNII: 9E8X80D2L0)		
Water (UNII: 059QF0KO0R)		

I	Packaging				
-	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:55505-222- 29	1 in 1 CARTON	01/01/2024		
	L	60 mL in 1 BOTTLE; 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	01/01/2024	

Labeler - Kramer Laboratories (122720675)

Revised: 1/2024 Kramer Laboratories