

NIZORAL SCALP ITCH RELIEF- hydrocortisone shampoo

Kramer Laboratories

Drug Facts

Active ingredient	Purpose
Hydrocortisone 1.0%	Anti-itch 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
LABORATORIES

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 Moiety

Ingredient Name	Basis of Strength	Strength
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Hydrocortisone (UNII: W4X0X7BPJ) (Hydrocortisone - UNII:W4X0X7BPJ)		Hydrocortisone	10 mg in 1 mL	
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: 6OZP39ZG8H)				
.Alpha.-Tocopherol Acetate (UNII: 9E8X80D2L0)				
Water (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55505-222-29	1 in 1 CARTON	01/01/2024	
1		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: 12/2025

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