HYDROCORTISONE MAXIMUM STRENGTH- hydrocortisone ointment Chain Drug Consortium, LLC

Hydrocortisone Ointment Maximum Strength - Premier Value

Drug Facts

Active ingredient

Hydrocortisone, USP 1%

Purpose

Anti-itch

Uses

- temporarily relieves itching associated with minor irritations, inflammation, and rashes due to
 - eczema
 - psoriasis
 - insect bites
 - o poison ivy, oak, sumac
 - detergents,
 - jewelry,
 - cosmetics
 - soaps
 - seborrheic dermatitis
- temporarily relieves external anal and genital itching
- other uses of this product should be only 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 a diaper rash. Consult.

When using this product

- avoid contact with the eyes
- do not use more than directed unless directed 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
- do not begin the use of any other hydrocortisone product unless directed by a doctor
- rectal bleeding occurs

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- for itching of skin irritation, inflammation, and rashes
 - adults and children 2 years of age and older: apply to the affected area not more than 3 to 4 times daily
 - o children under 2 years of age: consult a doctor
- for external anal and genital itching
 - adults: when practical, clean the affected area with mild soap and warm water; rinse thoroughly
 - gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product
 - apply to the affected area not more than 3 to 4 times daily
 - children under 12 years of age: consult a doctor

other information

- store at 15°-30°C (59°-86°F). Protect from freezing.
- before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredients

mineral oil, white petrolatum

Questions or comments?

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

Principal display panel

COMPARE TO THE ACTIVE INGREDIENT IN MAXIMUM STRENGTH CORTIZONE 10®*

MAXIMUM STRENGTH

Hydrocortisone Ointment, USP 1%

ANTI-ITCH OINTMENT

Relieves

Itches and

Rashes

NET WT. OZ (g)

For the temporary relief of itching associated with minor skin irritations, inflammation and rashes

*This product is not manufactured or distributed by Chattem, Inc., distributor of Maximum Strength Cortizone 10 \$.

Distributed by:

Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

Package label



MAXIMUM STRENGTH

Hydrocortisone Ointment, USP 1%

ANTI-ITCH OINTMENT

Relieves Itches and Rashes



For the temporary relief of itching associated with minor skin irritations, inflammation and rashes

Premier Value

COMPARE TO THE ACTIVE INGREDIENT IN MAXIMUM STRENGTH CORTIZONE 108*

MAXIMUM STRENGTH Hydrocortisone Ointment, USP 1%

ANTI-ITCH OINTMENT

NET WT. 1 OZ (28 g)



Distributed by: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087



If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

Product of India



Code: GO/DRUGS/361 ADD3A/02 PLD-B699B FC007028

"This product is not manufactured or distributed by Chattem, Inc., distributor of Maximum Strength Contzone 10".

Questions or comments? Call 1-877-753-3936 Monday-Friday 9AM-5PM EST

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PREMIER VALUE Maximum Strength Anti-Itch Ointment

HYDROCORTISONE MAXIMUM STRENGTH

hydrocortisone ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-009
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		
MINERAL OIL (UNII: T5L8T28FGP)			
PETROLATUM (UNII: 4T6H12BN9U)			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68016-009- 01	1 in 1 CARTON	09/08/2006		
1		28 g in 1 TUBE; Type 0: Not a Combination Product			

Marketing Information				
Marketing	Application Number or Monograph	Marketing Start	Marketing End	

Category	Citation	Date	Date
OTC Monograph Drug	M017	09/08/2006	

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 10/2023 Chain Drug Consortium, LLC