AQUAPHOR ITCH RELIEF- hydrocortisone ointment Beiersdorf Inc

Aquaphor Itch Relief Ointment

Active Ingredient

Hydrocortisone 1%

Purpose

Anti-Itch Ointment

Uses

- for the temporary relief of itching associated with minor skin irritations, inflammation, and rashes due to:
- eczema
- insect bites
- soaps
- poison ivy, oak, sumac
- seborrheic dermatitis
- psoriasis
- detergents
- cosmetics
- jewelry

Warnings

For external use only

Do not use

• for the treatment of diaper rash. Consult a doctor.

When using this product

- avoid contact with eyes
- do not exceed the recommended daily dosage unless directed by a doctor

Stop use and consult a doctor if

• condition worsens, if 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

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

Directions

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, consult a doctor.

Inactive ingredients

Petrolatum, Mineral Oil, Ceresin, Lanolin Alcohol,

Panthenol, Glycerin, Bisabolol, Menthoxypropanediol

Questions or comments?

1-800-227-4703

Aquaphor Itch Relief Ointment

Maximum Strength

Immediately Soothes & Relieves Itch & Irritiation

Up to 12 hours Significant Itch Relief

Clinically Proven

1% Hydrocortisone Anti-Itch Ointment

Dermatologist Recommended Brand

Fragrance free and Paraben free

Hypoallergenic

Helps Heal the Itch

Clinically proven significant itch relief for up to 12 hours

Immediately soothes itchy spots to help skin heal

Skin irritation

Skin rashes

Eczema

Psoriasis

Eczema

Poison Ivy



Children's

Aquaphor Itch Relief Ointment

Ages 2 Years and Older

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Number 1 Pediatrician Recommended Brand for Eczema

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AQUAPHOR ITCH RELIEF

hydrocortisone ointment

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:10356-120

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			
3-((L-MENTHYL)OXY)PROPANE-1,2-DIOL (UNII: KD6TZ2QICH)				
MINERAL OIL (UNII: T5L8T28FGP)				
CERESIN (UNII: Q1LS2UJO3A)				
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)				
GLYCERIN (UNII: PDC6A3C0OX)				
PANTHENOL (UNII: WW9CM0O67Z)				
LEVOMENOL (UNII: 24WE03BX2T)				
PETROLATUM (UNII: 4T6H12BN9U)				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:10356-120- 30	0.9 g in 1 PACKET; Type 0: Not a Combination Product	11/01/2020				
2	NDC:10356-120- 52	1 in 1 CARTON	11/01/2020				
2	NDC:10356-120- 04	28 g in 1 TUBE; Type 0: Not a Combination Product					
3	NDC:10356-120- 04	28 g in 1 TUBE; Type 0: Not a Combination Product	11/01/2020				
4	NDC:10356-120- 53	2 in 1 CARTON	11/01/2020				
4	NDC:10356-120- 04	28 g in 1 TUBE; Type 0: Not a Combination Product					
5	NDC:10356-120- 54	1 in 1 CARTON	06/01/2021				
5	NDC:10356-120- 07	56 g in 1 TUBE; Type 0: Not a Combination Product					
6	NDC:10356-120- 07	56 g in 1 TUBE; Type 0: Not a Combination Product	06/01/2021				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M017	11/01/2020			

Revised: 12/2023 Beiersdorf Inc