DERMFREE ANTI-ITCH- hydrocortisone1%, anti-itch cream
Jiangxi Hemei Pharmaceutical Co., Ltd

84010-021

Active Ingredient

Hydrocortisone 1%

Purpose

Anti-itch

Use

temporarily relieves pain and itching associated with:insect bites minor burns sunburn minor skin irritations minor cuts scrapes rashes due to poison ivy, poison oak, and poison sumac dries the oozing and weeping of poison ivy, poison oak, and poison sumac

Warnings

For external use only.

Do not use

on large areas of the body with any other product containing diphenhydramine, even one taken bymouth

When Using

avoid contact with eyes

Stop Use

condition worsens or does not improve within 15 dayssymptoms persist for more than 15 days or clear up and occur again withina few days

Ask Doctor

on chicken pox on measles

Keep Oot Of Reach Of Children

lf swallowed, get medical help or contact aPoison control center right away.

Directions

do not use more than directed 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: ask a doctor

Other information

protect from excessive heat (40°C/104°F)

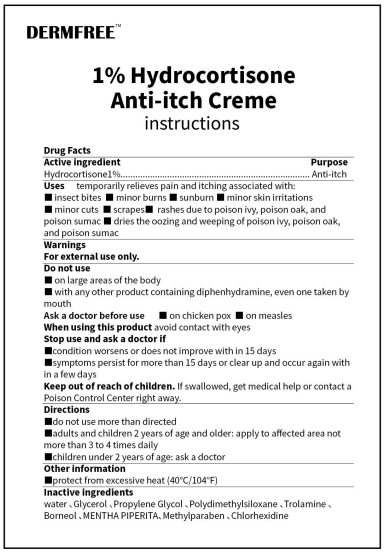
Inactive ingredients

water ，Glycerol ，Propylene Glycol ，Polydimethylsiloxane ，Trolamine，Borneol ，MENTHA PIPERITA,Methylparaben ,Chlorhexidine

PRINCIPAL DISPLAY PANEL

130*35

90*130



150*30*45

DERMFREE ANTI-ITCH

hydrocortisone1%, anti-itch cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84010-021
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**
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CHLORHEXIDINE (UNII: R4KO0DY52L)	
TROLAMINE (UNII: 9O3K93S3TK)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
BORNEOL (UNII: M89NIB437X)	
MENTHA PIPERITA (UNII: 79M2M2UDA9)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:84010-021-01	42.5 g in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/15/2024	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-021)

Revised: 8/2024 Jiangxi Hemei Pharmaceutical Co., Ltd