DERMFREE ANTI-ITCH- hydrocortisone acetate1%, anti-itch cream
Jiangxi Hemei Pharmaceutical Co., Ltd

84010-102

Active Ingredient

Hydrocortisone acetate 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

Stop Use

■condition worsens or does not improve within 15 days

■symptoms 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

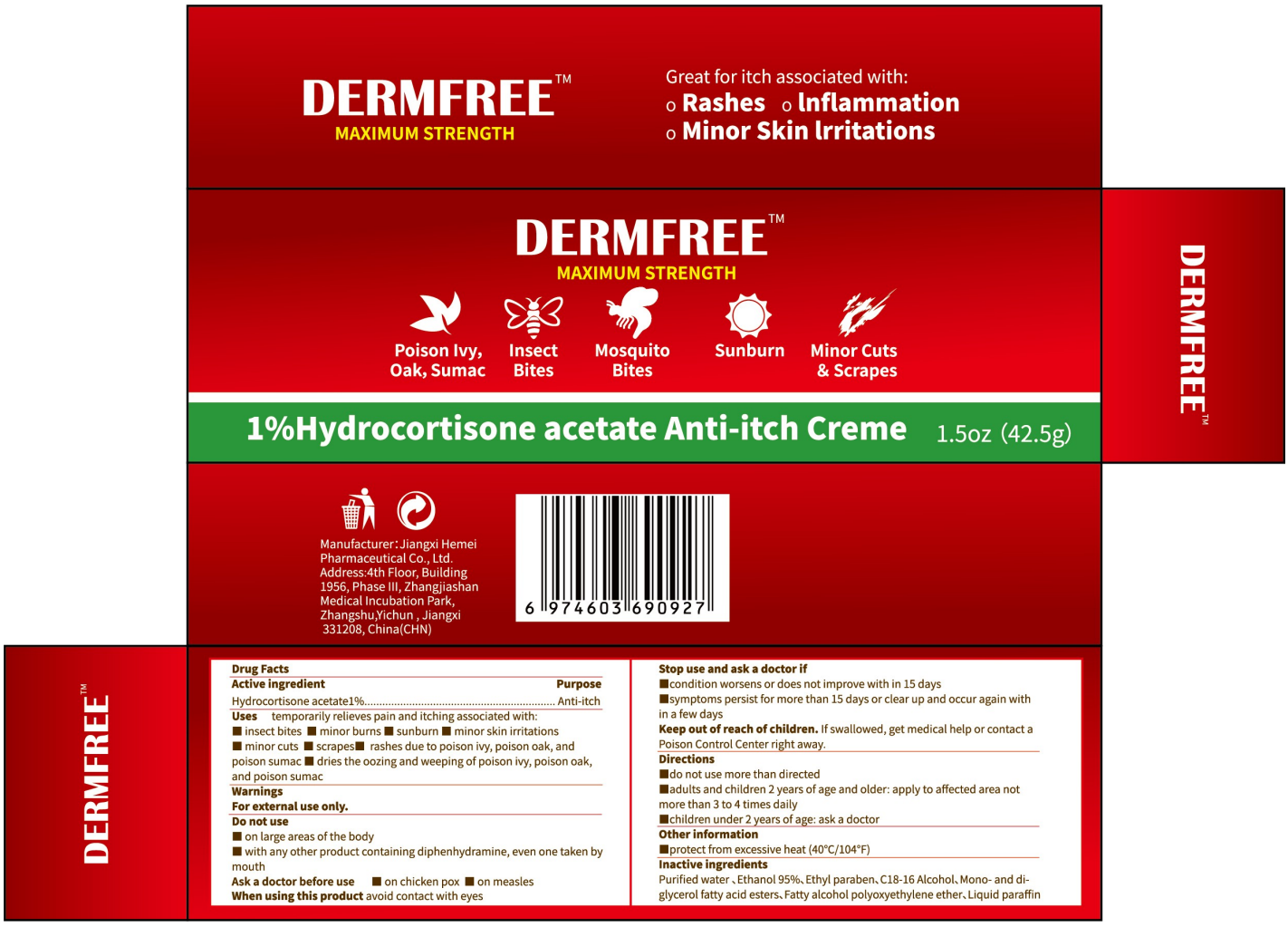
■protect from excessive heat (40°C/104°F)

Inactive ingredients

Purified water、Ethanol 95%、Ethyl paraben、C18-16 Alcohol、Mono- and di-glycerol fatty acid esters、Fatty alcohol polyoxyethylene ether、Liquid paraffin

PRINCIPAL DISPLAY PANEL

150*30*45



DERMFREE ANTI-ITCH
hydrocortisone acetate1%, anti-itch cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84010-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
LAURETH-9 (UNII: 0AWH8BFG9A)	
ETHYLPARABEN (UNII: 14255EXE39)	
PARAFFINUM LIQUIDUM (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-102-01	42.5 g in 1 BOTTLE; Type 0: Not a Combination Product	04/22/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/22/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-102)

Revised: 4/2025 Jiangxi Hemei Pharmaceutical Co., Ltd