DERMFREE ANTI-ITCH CREAM- hydrocortisone1% anti-itch cream cream
Jiangxi Zhongletang Pharmaceuticals Co.,Ltd.

87686-001

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 by mouth

When Using

avoid contact with eyes

Stop Use

■condition worsens or does not improve within 15 days

■symptoms persist for more than 15 days or clear up and occur again within a few days

Ask Doctor

■condition worsens or does not improve within 15 days

■symptoms persist for more than 15 days or clear up and occur again within a few days

Keep Out Of Reach Of Children

If swallowed, get medical help or contact a Poison 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, Polysorbate 80, Phenoxyethanol, C18-16 Alcohol, Glyceryl Stearate, Stearic Acid, Liquid Paraffin, 95%(v/v) Ethanol

PRINCIPAL DISPLAY PANEL

DERMFREE ANTI-ITCH CREAM

hydrocortisone1% anti-itch cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87686-001
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
STEARIC ACID (UNII: 4ELV7Z65AP)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
PARAFFINUM LIQUIDUM (UNII: T5L8T28FGP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87686-001-01	42.5 g in 1 TUBE; Type 0: Not a Combination Product	05/12/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/12/2026	

Labeler - Jiangxi Zhongletang Pharmaceuticals Co.,Ltd. (411295811)

Revised: 5/2026 Jiangxi Zhongletang Pharmaceuticals Co.,Ltd.