

FLOCEAN ANTI-ITCH- benzocaine 5%anti-itch cream
Jiangxi Hemei Pharmaceutical Co., Ltd

84010-017

Active Ingredient

Benzocaine 5%

Purpose

External analgesic

Use

temporarily relieves itching

Warnings

For external use only

Do not use

if you have a vaginal discharge. Consult a physician.

When Using

Avoid contact with eyes

Stop Use

condition worsens, or if symptoms persist formore than 7 days, or clear up and occur again within a few days Do not applyover large areas of the body

Ask Doctor

condition worsens, or if symptoms persist formore than 7 days, or clear up and occur again within a few days Do not applyover large areas of the body

Keep Oot Of Reach Of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 12 years and older apply a fingertip amount (approximately 1-inch strip) to affected area not more than 3 to 4 times daily, Clean nozzle of tube by wiping thoroughly before replacing cap. Keep cap tightly closed between uses.

Other information

children under 12 years consult a doctor

Inactive ingredients

water ,Glycerol ,Propylene Glycol ,Polydimethylsiloxane ,Trolamine,Borneol,MENTHAPIPERITA,Methylparaben ,Chlorhexidine

PRINCIPAL DISPLAY PANEL



FLOCEAN ANTI-ITCH

benzocaine 5% anti-itch cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:84010-017

Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)		BENZOCAINE	5 g in 100 g	
Inactive Ingredients				
Ingredient Name			Strength	
METHYLPARABEN (UNII: A2I8C7HI9T)				
DIMETHICONE (UNII: 92RU3N3Y1O)				
TROLAMINE (UNII: 9O3K93S3TK)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
BORNEOL (UNII: M89NIB437X)				
MENTHA PIPERITA (UNII: 79M2M2UDA9)				
CHLORHEXIDINE (UNII: R4KO0DY52L)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-017-01	28 g in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2024	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017		06/25/2024	

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	label(84010-017) , manufacture(84010-017)	

Revised: 6/2024

Jiangxi Hemei Pharmaceutical Co., Ltd