

999 ITCH RELIEVING- hydrocortisone ointment
Guangdong CR. Shunfeng Pharmaceutical Co Ltd

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

999 ITCH RELIEVING OINTMENT

ACTIVE INGREDIENTS

Hydrocortisone 1.0%

Purpose

Anti-Itch

Use

For the temporary relief of itching associated with minor skin irritations, inflammation and rashes due to: cosmetics, insect bites, psoriasis, eczema, seborrheic dermatitis.

Warnings

For external use only. Avoid contact with the eyes. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and seek medical advice if

Condition worsens. Symptoms persist for more than 7 days. Symptoms clear up and occur again within a few days. Excessive irritation of the skin develops.

Directions

Adult and children 2 years of age and older. Apply liberally to affected area not more than 5 times daily and 5-8 times per day for some severe cases or follow doctor's instruction.

Pregnant and children under 2 years of age. Do not use, consult a doctor.

Keep out of reach of children.

Other Information

Keep in a tightly closed container. Store at 8 to 30 ° (46-86°F) in a dry place away from sunlight.

You may report serious side effects to: 1452 W. Holt Ave, Pomona, CA 91768

Inactive Ingredients

Glyceryl Monostearate

Glyceryl Distearate

Glycerin
 Stearic Acid
 Mineral Oil
 Petrolatum
 Paraffin
 Ceresin
 Beta Carotene
 Ethylparaben
 Sodium Lauryl Sulfate

Drug Facts



文件名	A01003A6 皮炎平软膏纸盒[美国麦斯特201401版]彩样9. ai				
颜色	C M Y K 专银	尺寸	40*23*135mm 条码等级		
制作人	谢顺秀	日期	02-06 文件路径		\\serverctp\制作区\YXH苑晓辉\2014年8月\CP2014-08-1250\out
请对此稿的内容进行确认: 文字、尺寸、颜色、版式			客户签字		

Manufactured By Guangdong CR. Shunfeng Pharmaceutical Co.; Ltd.
 Jinjuzu, Dalinag Street, Shunde District, Foshan, Guangdong, China 528300

Distributed by Master Herbs Inc.
 1452 W. Holt Ave, Pomona, CA 91768
 (866)487-8373

999 ITCH RELIEVING

hydrocortisone ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69436-930
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCERYL DISTEARATE (UNII: 73071MW2KM)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
PARAFFIN (UNII: I9O0E3H2ZE)	
CERESIN (UNII: Q1LS2UJO3A)	
BETA CAROTENE (UNII: 01YAE03M7J)	
ETHYL PARABEN (UNII: 14255EXE39)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69436-930-20	1 in 1 BOX		
1		20 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/20/1989	

Labeler - Guangdong CR. Shunfeng Pharmaceutical Co Ltd (654033877)

Registrant - Guangdong CR. Shunfeng Pharmaceutical Co Ltd (654033877)

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong CR. Shunfeng Pharmaceutical Co Ltd		654033877	manufacture(69436-930)

Revised: 2/2015

Guangdong CR. Shunfeng Pharmaceutical Co Ltd