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.

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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□) 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, White Petrolatum, Ethylparaben, Sodium Lauryl Sulfate, Water

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

Drug Facts

999 ITCH RELIEVING
hydrocortisone ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69436-931
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)	
PETROLATUM (UNII: 4T6H12BN9U)	
ETHYLPARABEN (UNII: 14255EXE39)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69436-931-20	1 in 1 BOX	07/20/1989	
1		20 g in 1 TUBE; Type 0: Not a Combination Product		

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(69436-931)

Revised: 11/2016 Guangdong CR. Shunfeng Pharmaceutical Co Ltd