HEMORRODIL UNGUENTO PLUS- hydrocortisone ointment ZURICH MEDICAL LABS, LLC

Hemorrodil Unguento Plus

Drug Facts

Active Ingredients

Hydrocortisone (1%)

Purpose

Anti-Itch

Uses

Temporary relief of external anal itch & minor skin irritations and rashes.

Warnings

For external use only.

Do not use for treatment of diaper rash. Consult a doctor.

When using this product: avoid contact with eyes, do not exceed the recommended daily dosage unless directed by a doctor, & do not put into the rectum by using fingers or any mechanical device or applicator.

Stop use and ask a doctor if: bleeding occurs, condition worsens, &/or symptoms persist for more than 7 days or clear up & occur again with a few days. Do not begin use of any other hydrocortisone product unless you have consulted a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Instructions

Adults

When practical, cleanse the affected area by patting or blotting with an appropriate cleansing wipe. Gently dry by patting or blotting with tissue or a soft cloth before application of this product

Adults and children 12 years of age and older: apply to the affected area not more than 3 to 4 times daily.

Children under 12 years of age

Do not use, consult a doctor.

Store at room temperature or in cool place, but not over 80°F.

Inactive Ingredients

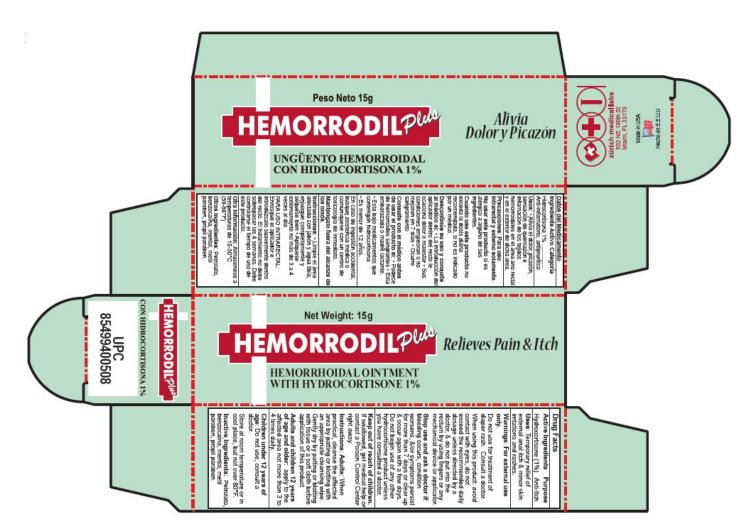
Petrolato, benzocaína, mentol, metil paraben, propil paraben.

PRINCIPAL DISPLAY PANEL - 15 g Tube Carton

Net Weight: 15g

HEMORRODIL Plus Relieves Pain & Itch

HEMORRHOIDAL OINTMENT WITH HYDROCORTISONE 1%



HEMORRODIL UNGU	JENTO PLUS					
hydrocortisone ointment						
Product Information						
Product T ype	HUMAN OTC DRUG	Item Code (Sou	Item Code (Source)		NDC:61357-132	
Route of Administration	RECTAL					
Active Ingredient/Active M	Ioiety					
Ingredient Name			Basis of Strength		Strength	
Hydrocortisone (UNII: WI4X0X7BPJ) (Hydrocortisone - UNII:WI4X0X7BPJ)			Hydro cortiso ne		10 mg in 1 g	

Ingredient Name					Strength	
BENZOCAINE (UNII: U3F	RS Y48 JW5)					
PETROLATUM (UNII: 4T	6H12BN9U)					
MENTHOL (UNII: L7T10	EIP3A)					
METHYLPARABEN (UNI	I: A218C7HI9T)					
PROPYLPARABEN (UNI	I: Z8IX2SC1OH)					
Packaging						
00	Package Description	Marketing	g Start Date	Ma	rketing End Date	
# Item Code	Package Description 1 in 1 CARTON	Marketing	g Start Date	Ma	rketing End Date	
Item Code NDC:61357-132-01	v	Marketin	g Start Date	Ma	rketing End Date	
 Packaging Item Code NDC:61357-132-01 1 	1 in 1 CARTON	Marketing	g Start Date	Ma	rketing End Date	
Item Code NDC:61357-132-01	1 in 1 CARTON	Marketin	g Start Date	Ma	rketing End Date	
Item Code 1 NDC:61357-132-01 1	1 in 1 CARTON 15 g in 1 TUBE	Marketin	g Start Date	Ma	rketing End Date	
Item Code NDC:61357-132-01	1 in 1 CARTON 15 g in 1 TUBE		g Start Date Marketing Start E		rketing End Date Marketing End Date	

Labeler - ZURICH MEDICAL LABS, LLC (071904097)

Establishment

Name	Address	ID/FEI	Business Operations
ZURICH MEDICAL LABS, LLC		071904097	MANUFACTURE(61357-132)

Revised: 3/2014

ZURICH MEDICAL LABS, LLC