

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 UNGUENTO PLUS

hydrocortisone ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:6 1357-132
Route of Administration	RECTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Hydrocortisone (UNII: W14X0 X7BPJ) (Hydrocortisone - UNII:W14X0 X7BPJ)	Hydrocortisone	10 mg in 1g

Inactive Ingredients

Ingredient Name	Strength
BENZOCAINE (UNII: U3RSY48JW5)	
PETROLATUM (UNII: 4T6H12BN9U)	
MENTHOL (UNII: L7T10EIP3A)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61357-132-01	1 in 1 CARTON		
1		15 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
EXPORT ONLY		03/01/1964	

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