LEADER MEDICATED- miconazole nitrate powder Cardinal Health Inc.

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.

Drug Facts

Active ingredient

Miconozole nitrate2%

Purpose

Antifungal

Uses

for the cure of most athlete's foot, jock itch and ringworm

Warnings

For external use only.

Do not use

on children under 2 years of age unless directed by a doctor

When using this product

do not get into eyes.

Stop use and ask a doctor if

if irritation occurs or there is no improvement within 4 weeks for athlete's foot and ringworm, or 2 weeks for jock itch.

Keep out of reach of children

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

Directions

- clean and dry affected area
- apply a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor
- supervise children in the use of this product
- for athlete's foot, pay special attention to spaces between toes, wear well-fitting, ventilated shoes and change shoes and socks at least once daily
- for athlete's foot and ringworm, use daily for 4 weeks; for jock itch, use daily for 2 weeks
- if condition persists, consult a doctor
- this product is not effective on scalp or nails

Other infomation

Product settles during shipment. Package contains full net weight.

Inactive ingredients

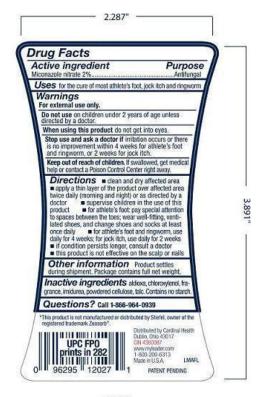
aldioxa, chloroxylenol, fragrance, imidurea, powdered cellulose, talc. Contains no starch

Questions

Call 1-866-964-0939

Box Label





BACK

LABEL SIZE 2.287"W x 3.891"H



Leader Medicated Miconazole Nitrate 2% Powder Spray

LEADER MEDICATED miconazole nitrate powder Product Information Product Type HUMAN OTC DRUG LABEL Item Code (Source) NDC:37205-653 Route of Administration TOPICAL DEA Schedule

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MICO NAZO LE NITRATE (MICONAZOLE)	MICONAZOLE	1.42 g in 71 g		

nactive Ingredients				
Ingredient Name	Strength			
ALDIO XA				
CHLOROXYLENOL				
IMIDUREA				
PO WDERED CELLULO SE				
TALC				

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:37205-653-18	71 g in 1 PACKAGE				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333C	10/31/2010				

Labeler - Cardinal Health Inc. (097537435)

Revised: 10/2012 Cardinal Health Inc.