

**INLIFAY ANTIFUNGAL- miconazole nitrate 2% antifungal cream**  
**Jiangxi Hemei Pharmaceutical Co., Ltd**

-----

**84010-252**

**Active Ingredient**

Miconazole Nitrate 2% w/w

**Purpose**

Antifungal

**Use**

Helps relieve the itching, burning, cracking, and scaling associated with fungal infections such as athlete's foot, jock itch, and ringworm.

**Warnings**

For external use only.

**Do not use**

you are allergic to miconazole or any of the ingredients in this product

**When Using**

Avoid contact with eyes. If contact occurs, rinse thoroughly with water.  
Do not use on children under 2 years of age unless directed by a doctor.

**Stop Use**

Irritation occurs Condition persists for more than 2 weeks or worsens

**Ask Doctor**

Irritation occurs Condition persists for more than 2 weeks or worsens

**Keep Out Of Reach Of Children**

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

**Directions**

Clean the affected area and dry thoroughly. Apply a thin layer to the affected area twice daily (morning and night). Supervise children in the use of this product. Use daily for the full treatment period as directed. This product is not effective on the scalp or nails.

Other information

1. Keep in a cool and dry place, avoid direct sunlight. 2. Keep out of the reach of children to avoid accidental ingestion.

Inactive ingredients

Aloe, Borneol, Cnidium, Dictamnus, Euphorbia, Glycerin, Light liquid paraffin, Medical petrolatum, Natural menthol, Purified water, Sophora Flavescens, Stearic acid, Stemona, Triethanolamine, Vitamin E

PRINCIPAL DISPLAY PANEL



INLIFAY ANTIFUNGAL			
miconazole nitrate 2% antifungal cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84010-252

Route of Administration	TOPICAL
-------------------------	---------

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII: 7NNO0D7S5M)	MICONAZOLE NITRATE	2 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	
WATER (UNII: 059QF0KO0R)	
STEMONA TUBEROSA ROOT (UNII: 7S9328671Z)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
ORIGANUM DICTAMNUS FLOWERING TOP (UNII: 9RBR98LSD9)	
BORNEOL (UNII: M89NIB437X)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
EUPHORBIA HIRTA (UNII: L13YF113GN)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	
CNIDIUM SEED (UNII: V1IA3S3CUS)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
ALOE (UNII: V5VD430YW9)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-252-01	100 g in 1 BOTTLE; Type 0: Not a Combination Product	01/22/2026	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	01/22/2026	

**Labeler** - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)**Establishment**

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-252)