DB ANTIFUNGAL FOOT- miconazole nitrate cream
The Magni Group Inc

DB Antifungal foot

Drug Facts

Active ingredients (in cream)

Miconazole Nitrate 2%

Purpose

Antifungal

Uses

• Cures most tinea pedis (athlete’s foot) • Relieves itching, scaling, cracking, burning, redness; itchy, scaly skin between toes; itching, burning feet; discomfort

Warnings

Do not use

- on children under 2 years of age unless directed by a doctor • For external use only
- Avoid contact with the eyes
- If irritation occurs or if there is no improvement within 4 weeks, discontinue use and consult a doctor

Keep out of reach of children

• If swallowed, get medical attention, help, or contact a poison control center right away

Directions

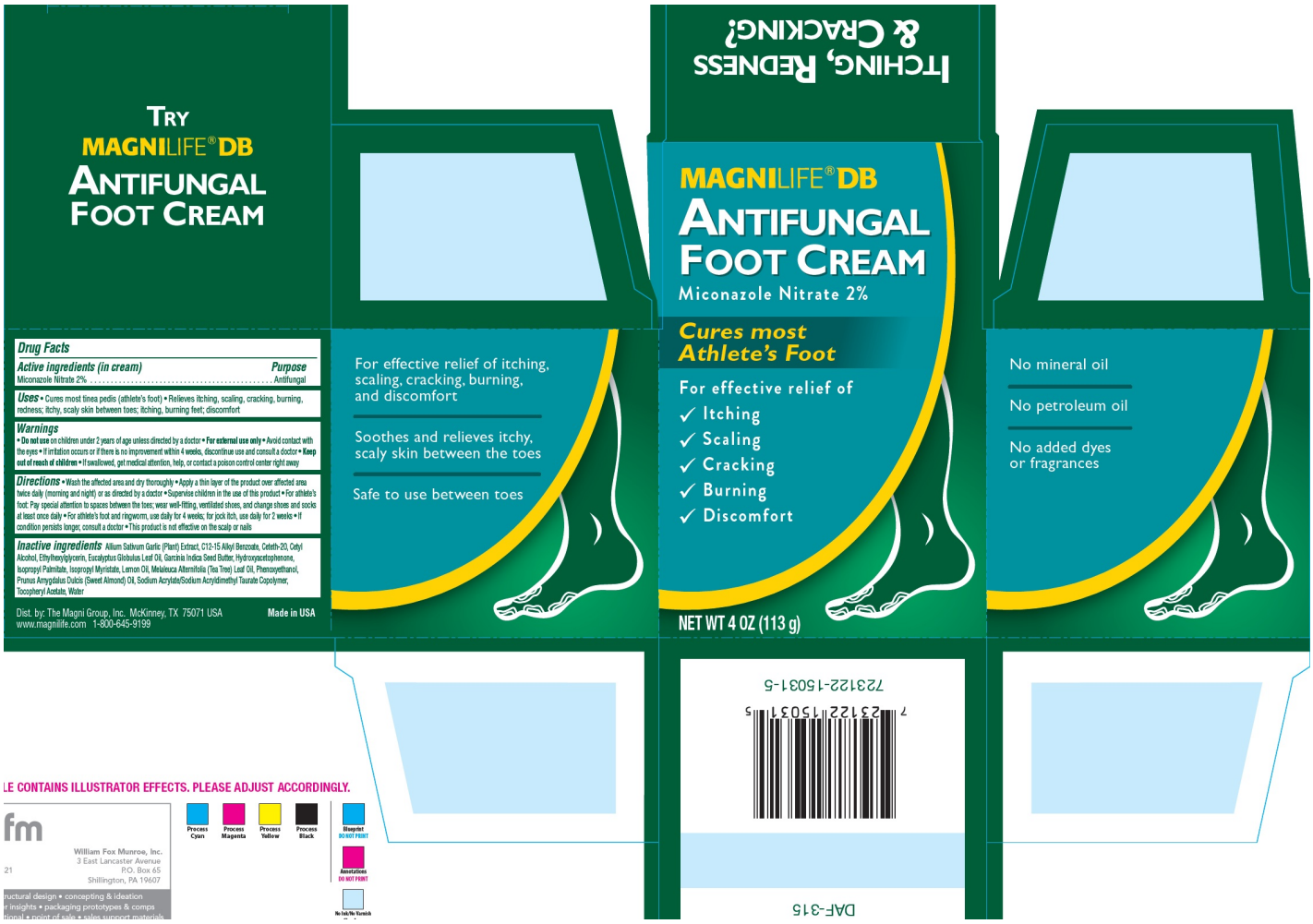
• Wash the affected area and dry thoroughly • Apply a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor • Supervise children in the use of this product • For athlete’s foot: Pay special attention to spaces between the toes; wear well-fitting, ventilated 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 longer, consult a doctor • This product is not effective on the scalp or nails

Inactive ingredients

Allium Sativum Garlic (Plant) Extract, C12-15 Alkyl Benzoate, Ceteth-20, Cetyl Alcohol, Ethylhexylglycerin, Eucalyptus Globulus Leaf Oil, Garcinia Indica Seed Butter,

Hydroxyacetophenone, Isopropyl Palmitate, Isopropyl Myristate, Lemon Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Phenoxyethanol, Prunus Amygdalus Dulcis (Sweet Almond) Oil, Sodium Acrylate/Sodium Acryldimethyl Taurate Copolymer, Tocopheryl Acetate, Water

Package Labeling:



DB ANTIFUNGAL FOOT

miconazole nitrate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43689-0025
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CETETH-20 (UNII: I835H2IHHX)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GARCINIA INDICA SEED BUTTER (UNII: US2H3D7800)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LEMON OIL (UNII: I9GRO824LL)	
TEA TREE OIL (UNII: VIF565UC2G)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALMOND OIL (UNII: 66YXD4DKO9)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43689-0025-1	113 g in 1 JAR; Type 0: Not a Combination Product	04/27/2017	
2	NDC:43689-0025-2	1 in 1 BOX	04/27/2017	
2		113 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	04/27/2017	

Labeler - The Magni Group Inc (113501902)

Revised: 10/2023 The Magni Group Inc