BENGAMA ANTIFUNGAL- tolnaftate solution
Genuine Drugs

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.

Bengama Antifungal Solution

Active ingredient (in each gram)

Tolnaftate 1%

Purpose

Antifungal

Uses

- clinically proven to cure most athlete's foot (tinea pedis) and ringworm (tinea corporis)
- helps prevnet most athlete's foot from recurring when used daily
- effectively soothes and relieves symptoms of athlete's foot, including itching, burning and cracking

Warnings

For external use only

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 4 weeks

Do not use

on children under 2 years of age except under the advice and supervision of a doctor.

Keep out of reach of children.

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

Directions

- shake well before using
- apply generously to affected areas and massage gently until liquid is absorbed into

the skin
- for adults and children over 12, rub well on the afffected area. repeat 3-4 times daily
- for children 12 years of age or younger, consult a doctor before use

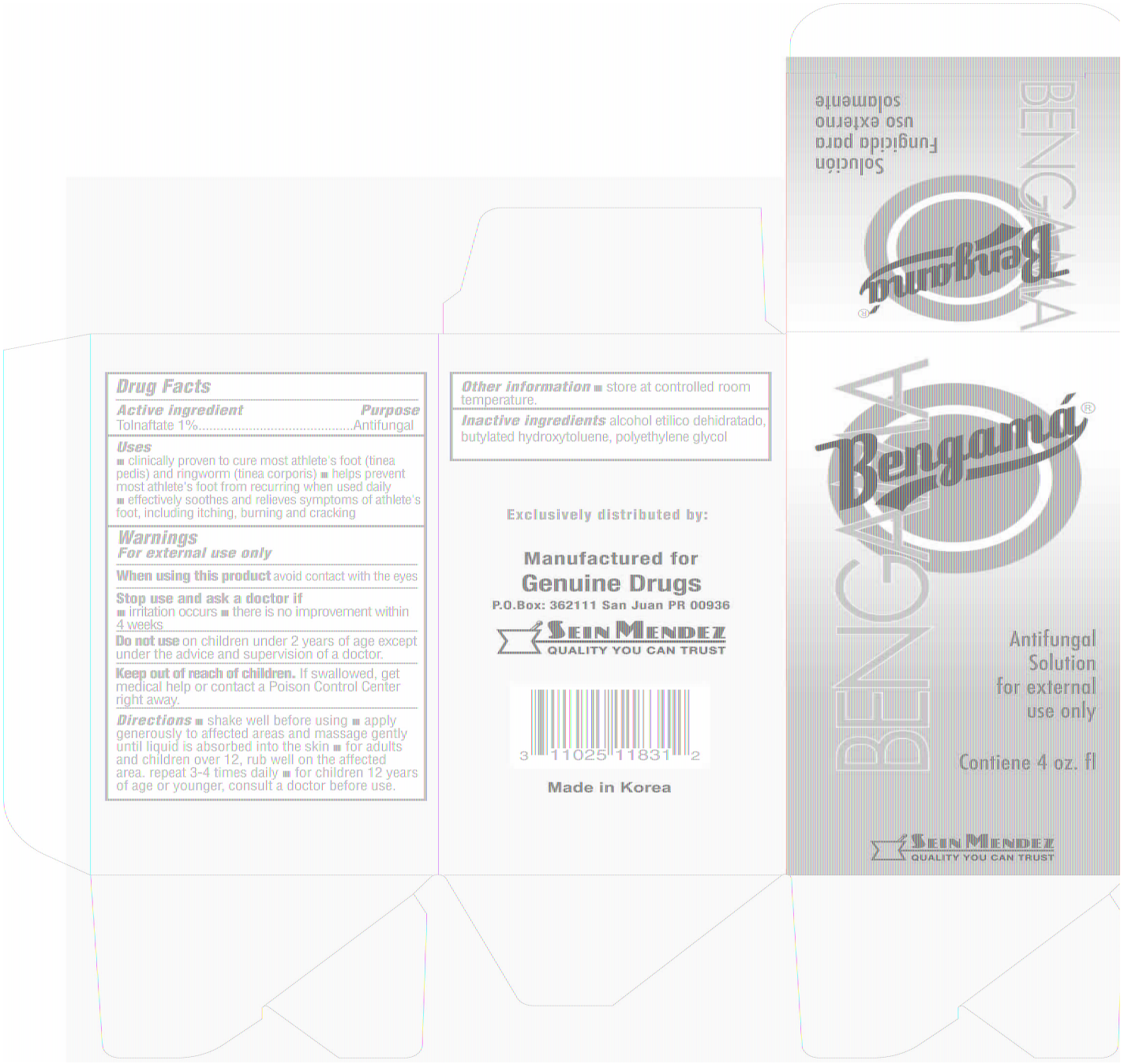
Inactive ingredients

dehydrated ethyl alcohol, butylated hydroxytoluene, polyethylene glycol

Other information
- store at controlled room temperature

Package label

Bengama antifungal



BENGAMA ANTIFUNGAL

tolnaftate solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69666-831
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	1 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69666-831-04	1 in 1 BOX	04/20/2015	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	04/20/2015	

Labeler - Genuine Drugs (079610378)

Revised: 2/2023 Genuine Drugs