MENTHOL- maximum strength medicated foot powder talc free powder
Topco Associates LLC

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.

Top Care Menthol Foot Powder Talc Free

Active ingredient

Menthol 1.0%

Purpose

External analgesic

Use

for the temporary relief of pain and itching associated with minor skin irritation

Warnings

For external use only.

When using this product
- avoid contact with eyes

Stop and consult a doctor if
- conditions worsens
- symptoms persists for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center right away.

Directions
- adults and children 2 years of age and older, apply to affected area not more than 3 to 4 times daily
- children under 2 years of age, consult a doctor
- wash and dry feet thoroughly
- sprinkle powder liberally on feet, between toes and on bottoms of feet

Inactive ingredients

benzathonium Chloride, eucalyptus oil, peppermint oil, sodium bicarbonate, tricalcium phosphate, zea mays (corn) starch

Questions?

Call - 1-888-423-0139

Principal Display Panel

Top Care health

MAXIMUM STRENGTH

Medicated Foot Powder

MENTHOL 1%

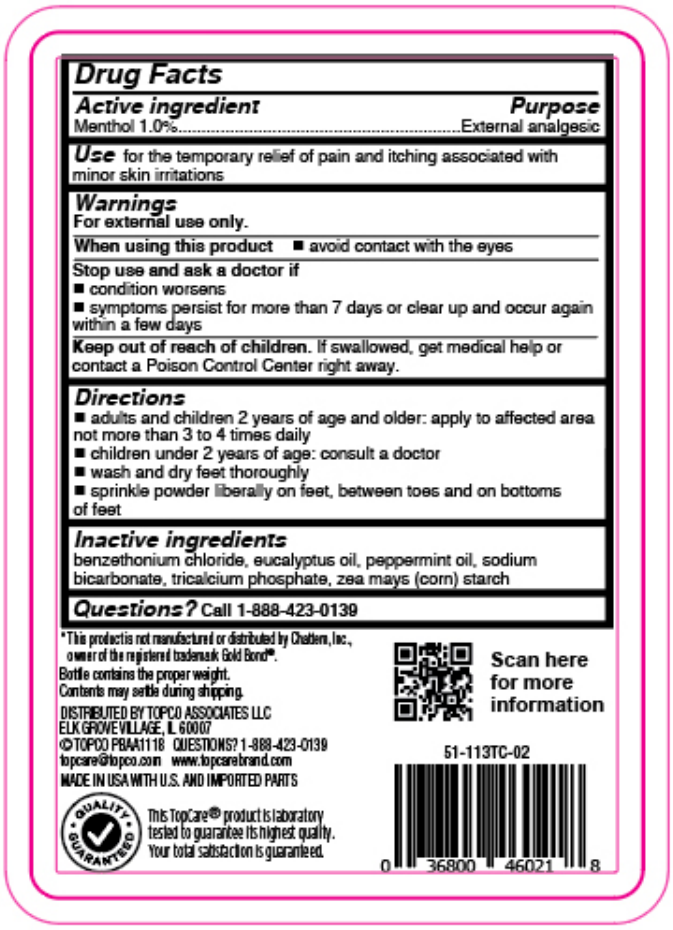
EXTERNAL ANALGESIC

TALC- FREE

Triple-Action Formula

- Absorbs Moisture
- Relieves Itching
- Helps Control Foot Odor

NET WT 10 OZ (283g)



MENTHOL

maximum strength medicated foot powder talc free powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:36800-252
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.8 g in 283 g

Inactive Ingredients	
Ingredient Name	**Strength**
ZEA MAYS SUBSP. MAYS WHOLE (UNII: 1G5HNE09V8)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-252-10	283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/10/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/10/2019	

Labeler - Topco Associates LLC (006935977)

Revised: 1/2023

Topco Associates LLC