MENTHOL- maximum strength medicated foot powder powder
VALU MERCHANDISERS, CO

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.

Medicated 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 irritations

Warnings

For external use only.

When using this product
- avoid contact with the eyes

Stop use and ask 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.

If swallowed, 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

benzethonium chloride, eucalyptus oil, peppermint oil, sodium bicarbonate, tricalcium phosphate, zea mays (corn) starch

Questions?

Call 1-866-964-0939

Principal Display Panel

Best Choice

MAXIMUM STRENGTH

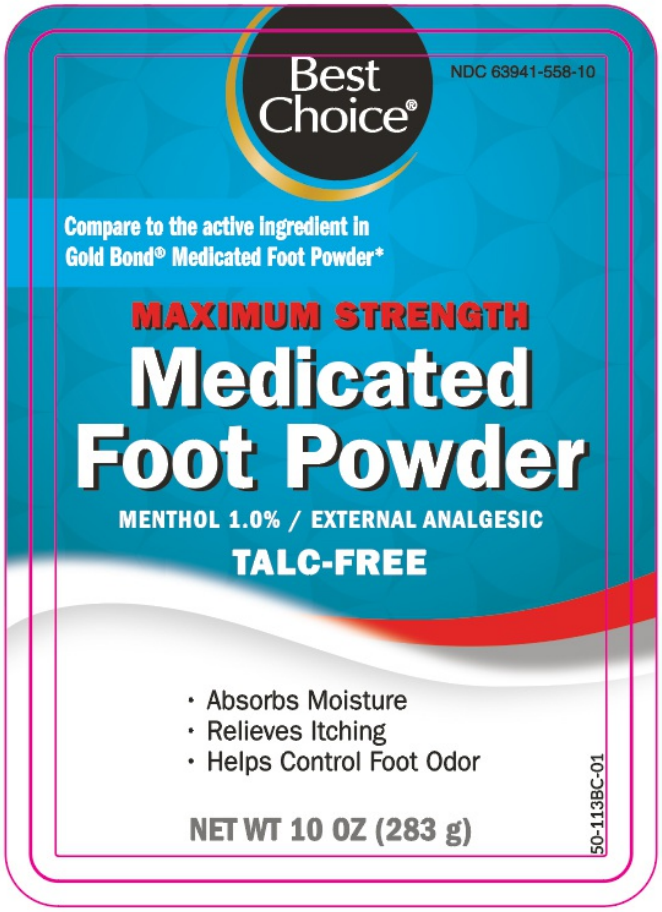
Medicated Foot Powder

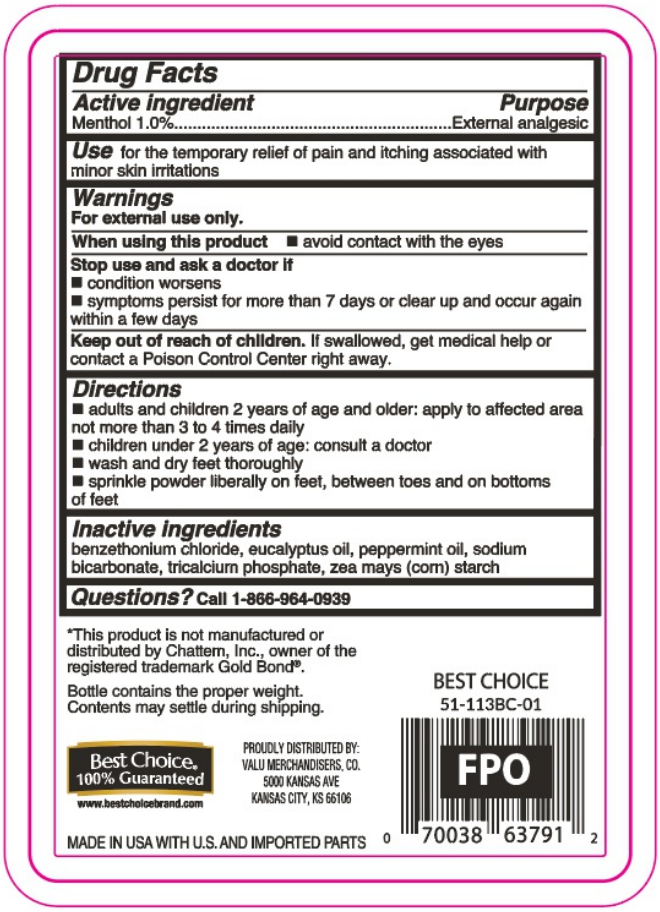
MENTHOL 1.0% / EXTERNAL ANALGESIC

TALC-FREE

- Absorbs Moisture
- Relieves Itching
- Helps Control Foot Odor

NET WT 10 oz (283g)





MENTHOL

maximum strength medicated foot powder powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63941-558
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**
EUCALYPTUS OIL (UNII: 2R04ONI662)	
STARCH, CORN (UNII: O8232NY3SJ)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63941-558-10	283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/15/2019	

Labeler - VALU MERCHANDISERS, CO (868703513)

Revised: 9/2019 VALU MERCHANDISERS, CO