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

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

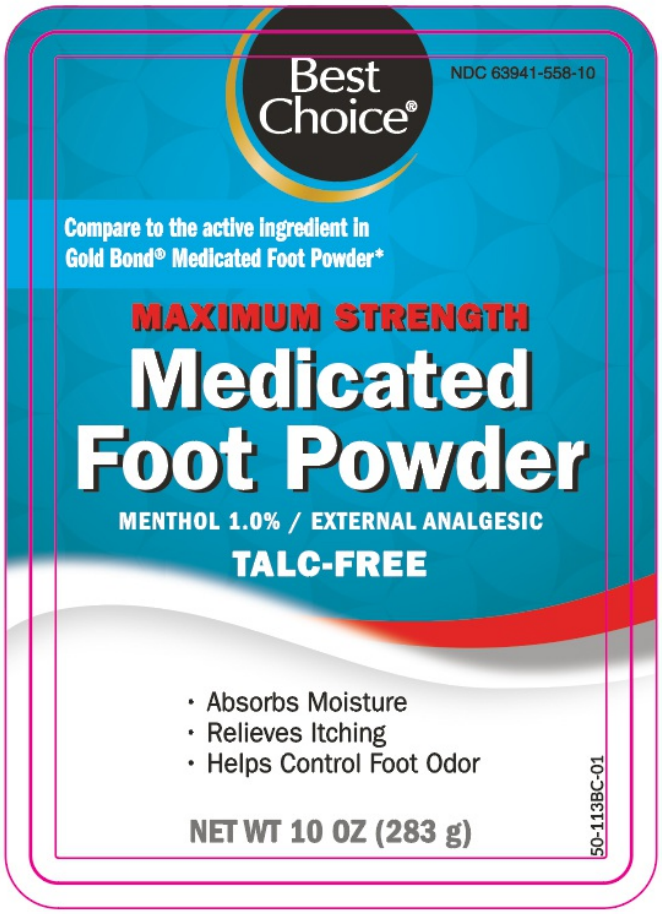
Medicated Foot Powder

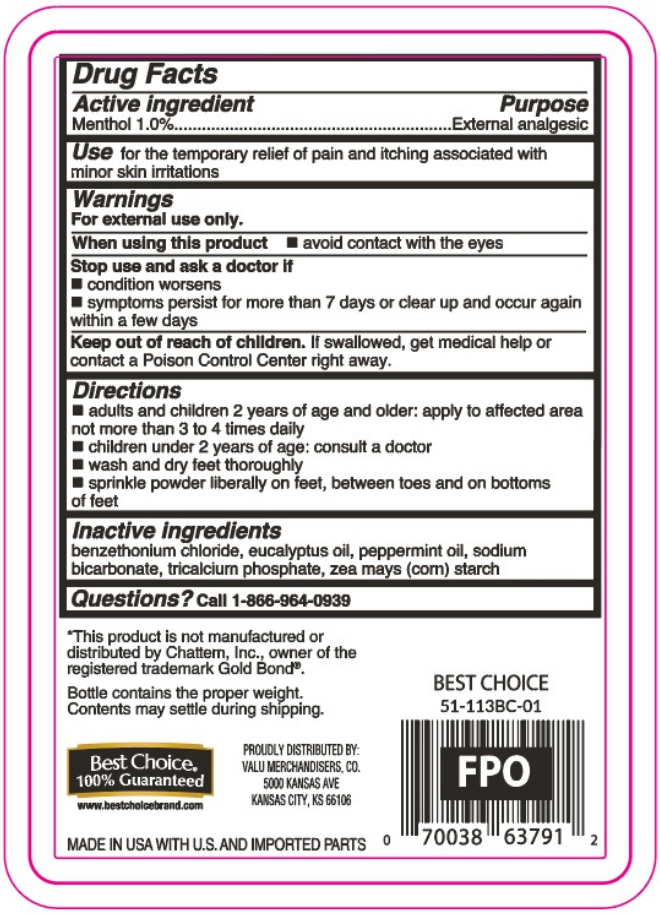
MENTHOL 1.0% / EXTERNAL ANALGESIC

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 Drug	M017	07/15/2019	

Labeler - VALU MERCHANDISERS, CO (868703513)

Revised: 2/2024 VALU MERCHANDISERS, CO