MENTHOL- maximum strength medicated foot powder talc free powder
Target Corporation

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 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 1-800-910-6874

Principal Display Panel

maximum strength

medicated

foot powder

menthol 1 %

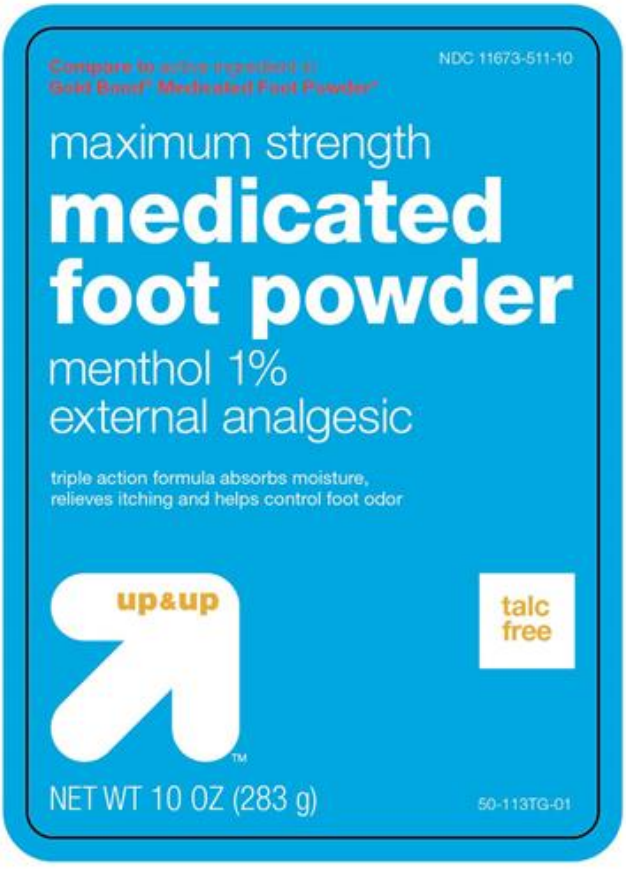
external analgesic

talc- free

triple action formula absorbs moisture,

relieve itching and helps control foot odor

NET WT 10 OZ (283 g)



MENTHOL
maximum strength medicated foot powder talc free powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-511
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)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-511-10	283 g in 1 BOTTLE, PLASTIC; Type 1: Convenience Kit of Co-Package	11/27/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/27/2017	

Labeler - Target Corporation (006961700)

Revised: 2/2024 Target Corporation