MAXIMUM STRENGTH MEDICATED FOOT POWDER- medicated foot powder powder
Target Corporation

Target Medicated Foot Powder Menthol 1%

Active ingredient

Menthol 1.0%

Purpose

External analgesic

Use

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

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

benzethonium chloride, eucalytus oil, gum acacia, peppermint oil, sodium bicarbonate,

talc

Questions?

call 1-800-910-6874

Principal Display Panel

Maximum strength

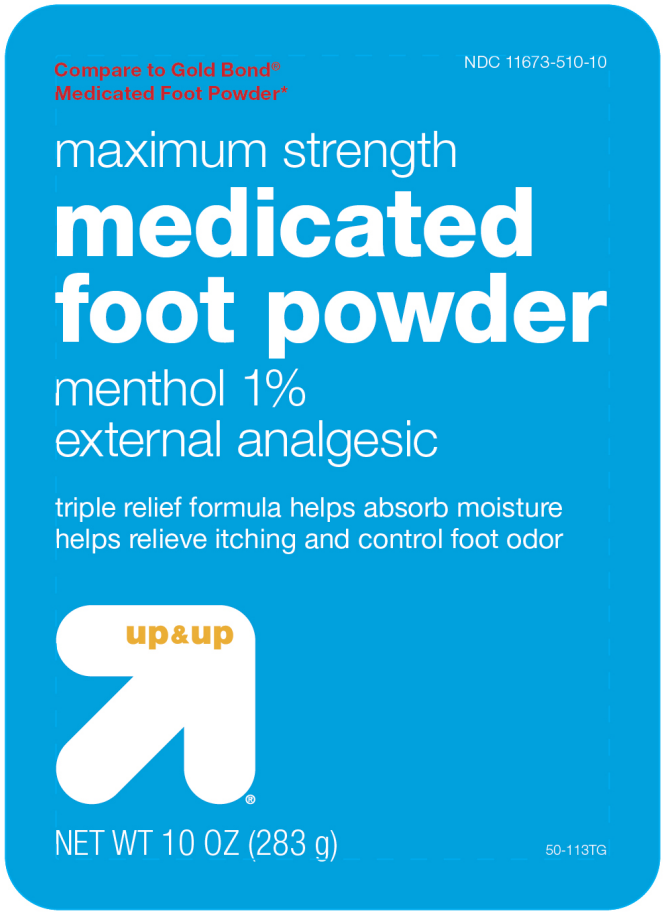
medicated

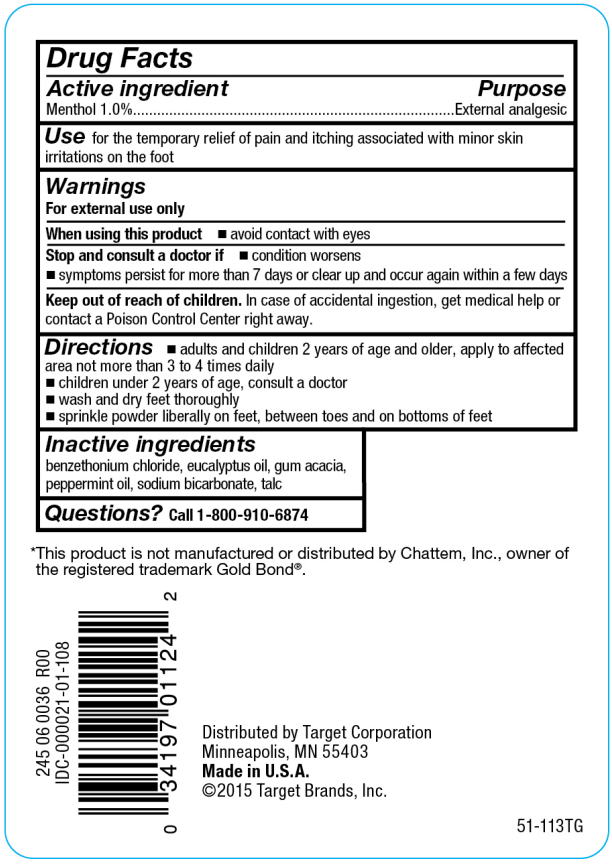
foot powder

menthol 1%
external analgesic

triple relief formula helps absorb moisture
helps relieve itching and control foot odor

NET WT 10 OZ (283g)





MAXIMUM STRENGTH MEDICATED FOOT POWDER

medicated foot powder powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-510
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.1 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
ACACIA (UNII: 5C5403N26O)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TALC (UNII: 7SEV7J4R1U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-510-10	283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/30/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/30/2013	

Labeler - Target Corporation (006961700)

Revised: 2/2024 Target Corporation