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

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?

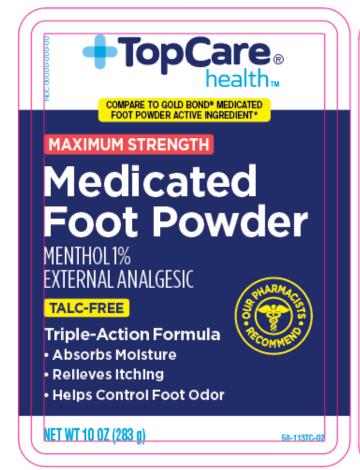
Call - 1-888-423-0139

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

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Prod	uct	ıntorn	nation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:36800-252

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)

MENTHOL

2.8 g in 283 g

Inactive Ingredients

Ingredient Name	Strength
(UNII: 1G5HNE09V8)	
4C08XP666)	

TRICALCIUM PHOSPHATE (UNII: K4C08XP666)

EUCALYPTUS OIL (UNII: 2R040NI662)

ZEA MAYS SUBSP. MAYS WHOLE

SODIUM BICARBONATE (UNII: 8MDF5V39QO)
BENZETHONIUM CHLORIDE (UNII: PH41D05744)

PEPPERMINT OIL (UNII: AV092KU4JH)

Packaging

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

Marketing Information

Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
M017	02/10/2019			
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date		

Labeler - Topco Associates LLC (006935977)

Revised: 12/2024 Topco Associates LLC