## GOLD BOND MEDICATED MAXIMUM STRENGTH FOOT- menthol powder Chattem, Inc.

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#### Gold Bond Medicated Maximum Strength Foot Powder - Reformulated

## **Gold Bond Medicated**

Foot Powder

**Drug Facts** 

## Active ingredient

## **Purpose**

#### Uses

temporarily relieves the pain and itch associated with:

minor skin irritations on the foot

## Warnings

#### For external use only

## When using this product

avoid contact with eyes

## Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- redness, irritation, swelling or pain persists or increases

#### Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

#### **Directions**

- **adults and children 2 years and older:** apply freely up to 3 or 4 times daily
- children under 2 years: ask a doctor
- thoroughly wash and dry feet, sprinkle powder liberally over feet, between toes and on bottoms of feet. Sprinkle inside shoes for maximum freshness.

## Inactive ingredients

zea mays (corn) starch, sodium bicarbonate, acacia senegal gum, silica, tricalcium phosphate, eucalyptus globulus leaf oil, mentha piperita (peppermint) oil, benzethonium chloride

## PRINCIPAL DISPLAY PANEL

TALC-FREE GOLD BOND MAXIMUM STRENGTH FOOT POWDER Net Wt 10 oz (283 g)





## **GOLD BOND MEDICATED MAXIMUM STRENGTH FOOT**

menthol powder

# Product Information Product Type HUMAN OTC DRUG Item Code (Source)

Ingradiant Name

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g	

NDC:41167-0171

## **Inactive Ingredients**

	ingredient wante	Strength
I	STARCH, CORN (UNII: O8232NY3SJ)	
	SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
	<b>ACACIA</b> (UNII: 5C5403N26O)	
	SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
l	TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
	FUCALYPTUS OII (UNII: 2R04ONI662)	

PEPPERMINT OIL (UNII: AV092KU4JH)
BENZETHONIUM CHLORIDE (UNII: PH41D05744)

## **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:41167- 0171-2	283 g in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2021	

	13 g in 1 BOTTLE; Type 0: Not a Combination roduct	03/01/2021		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	03/01/2021		

Labeler - Chattem, Inc. (003336013)

Revised: 10/2023 Chattem, Inc.