GOLD BOND MEDICATED MAXIMUM STRENGTH FOOT- menthol powder
Gold Bond Co LLC

Gold Bond Medicated Foot Powder

Gold Bond Medicated

Maximum Strength Foot Powder

Drug Facts

Active ingredient

Menthol 1%

Purpose

Anti-itch

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

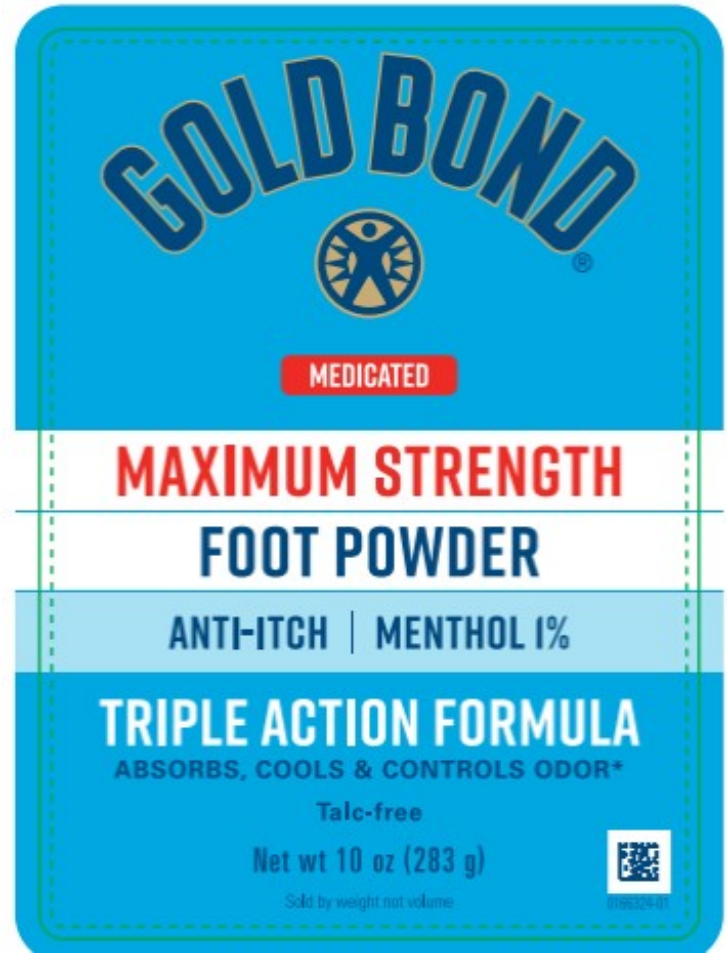
■ thoroughly wash and dry feet, sprinkle powder liberally over feet, between toes and on bottoms of feet. Sprinkle inside shoes for maximum freshness.

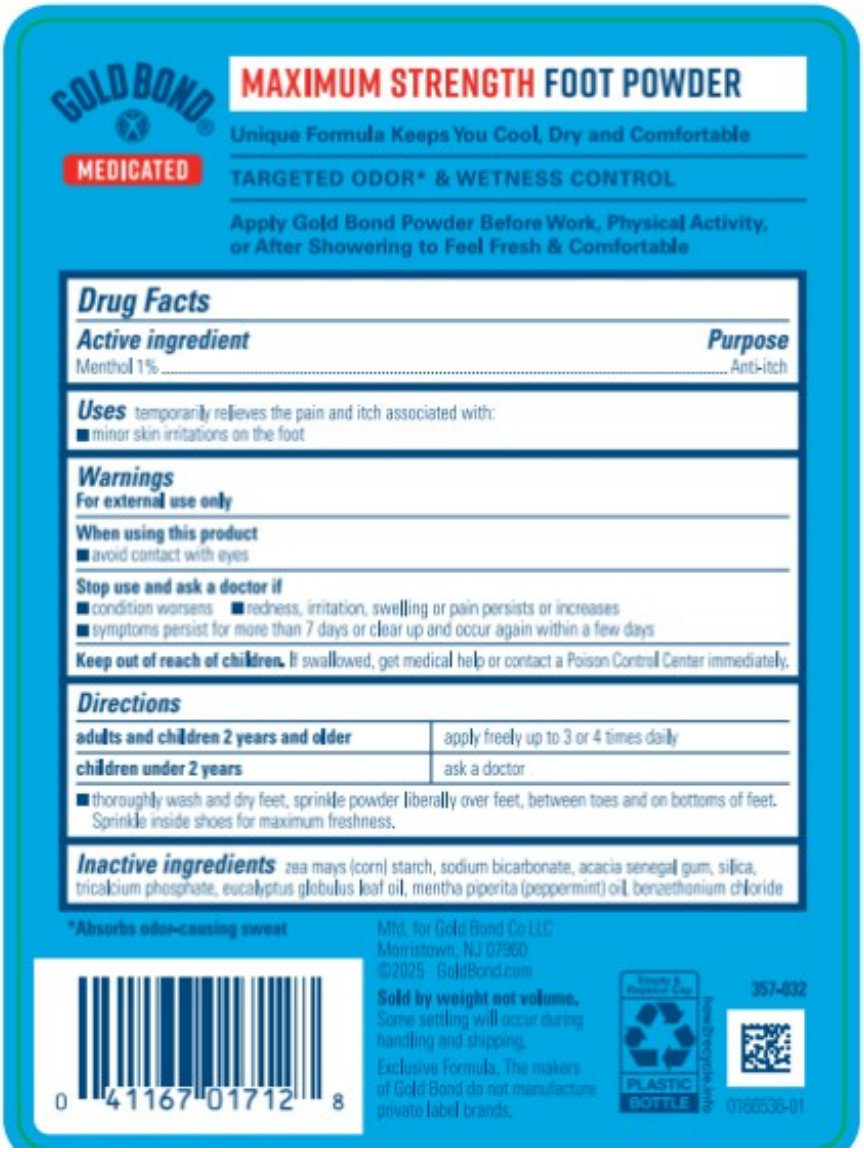
Inactive ingredients

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

GOLD BOND
MEDICATED
MAXIMUM STRENGTH
FOOT POWDER
ANTI-ITCH MENTHOL 1%
TRIPLE ACTION FORMULA
Net Wt 10 oz (283 g)





GOLD BOND MEDICATED MAXIMUM STRENGTH FOOT

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84714-0171
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

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	

ACACIA (UNII: 5C5403N26O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84714-0171-2	283 g in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2021	
2	NDC:84714-0171-6	113 g in 1 BOTTLE; Type 0: Not a Combination Product	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 - Gold Bond Co LLC (119184861)

Revised: 12/2025

Gold Bond Co LLC