

**GOLD BOND MEDICATED ORIGINAL STRENGTH BODY- menthol powder
Chattem, Inc.**

Gold Bond Medicated Original Strength Body Powder - Reformulated

**Gold Bond Medicated
Original Strength Body Powder
Drug Facts**

Active ingredient

Purpose

Menthol 0.15%.....
Anti-itch

Uses

temporarily relieves the pain and itch associated with:

- minor cuts ■ sunburn ■ insect bites ■ scrapes ■ minor burns ■ minor skin irritations

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

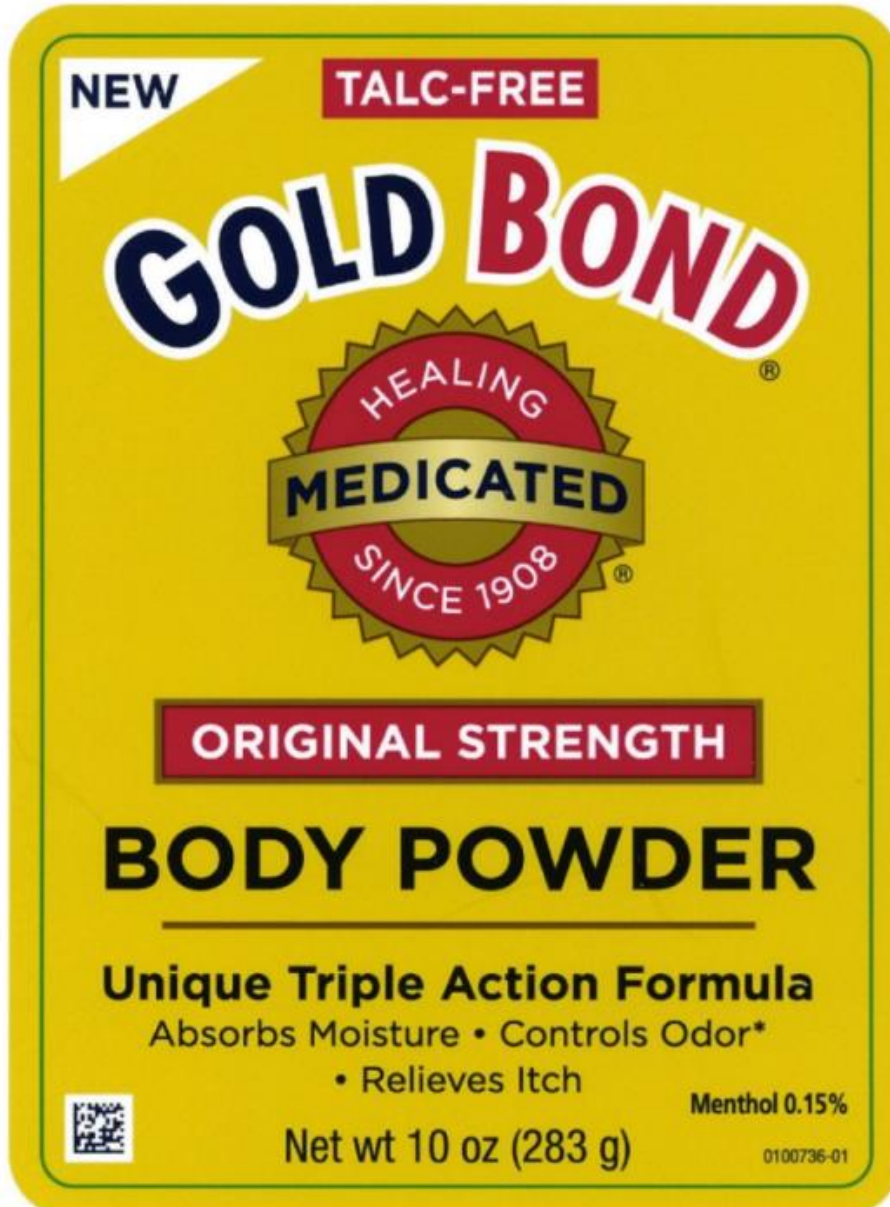
- **adults and children 2 years and older:** apply freely up to 3 or 4 times daily
- **children under 2 years:** ask a doctor
- for best results dry skin thoroughly before applying

Inactive ingredients

zea mays (corn) starch, zinc oxide, acacia senegal gum, silica, tricalcium phosphate, eucalyptol, methyl salicylate, salicylic acid, zinc stearate, thymol

PRINCIPAL DISPLAY PANEL

GOLD BOND
ORIGINAL STRENGTH
BODY POWDER
Net Wt 10 oz (283 g)



GOLD BOND® Body Powder Original Strength

Gold Bond® Medicated Original Strength Powder's unique triple action formula absorbs moisture, cools, and relieves itch so you feel fresh. It provides all over odor* and wetness control and contains a plant-based ingredient†. Apply Gold Bond before work, physical activity, or after showering to keep you cool, dry, and comfortable.

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Menthol 0.15%.....	Anti-itch

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*absorbs odor-causing sweat †refers to cornstarch 357-030
This product is sold by weight, not by volume. Some settling may occur.



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a Sanofi Company,
P. O. Box 2219, Chattanooga, TN
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www.GoldBond.com 0100737-01



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GOLD BOND MEDICATED ORIGINAL STRENGTH BODY

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41167-0110
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.15 g in 100 g	
Inactive Ingredients			
Ingredient Name	Strength		
STARCH, CORN (UNII: O8232NY3SJ)			
ZINC OXIDE (UNII: SOI2LOH54Z)			
ACACIA (UNII: 5C5403N26O)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			

TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
ZINC STEARATE (UNII: H92E6QA4FV)	
THYMOL (UNII: 3J50XA376E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-0110-9	283 g in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2021	
2	NDC:41167-0110-6	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 - Chattem, Inc. (003336013)

Revised: 10/2023

Chattem, Inc.