

TALC-FREE MEDICATED BODY- menthol powder
Davion, Inc

talc-free medicated body powder

Active Ingredient

Menthol 0.15%

Purpose

External Analgesic

Uses

Temporary relief of pain and itch associated with:

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

Warning

- **For external use only**
- 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 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

Remove seal beneath cap before first use. Do not use if seal is broken or missing.

- adults and children 2 years and older - apply freely upto 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, Tricalcium Phosphate, Sodium Bicarbonate, salicylic acid, zinc stearate, Eucalyptus oil, Peppermint Oil

PRINCIPAL DISPLAY PANEL

NDC 42669-209-01

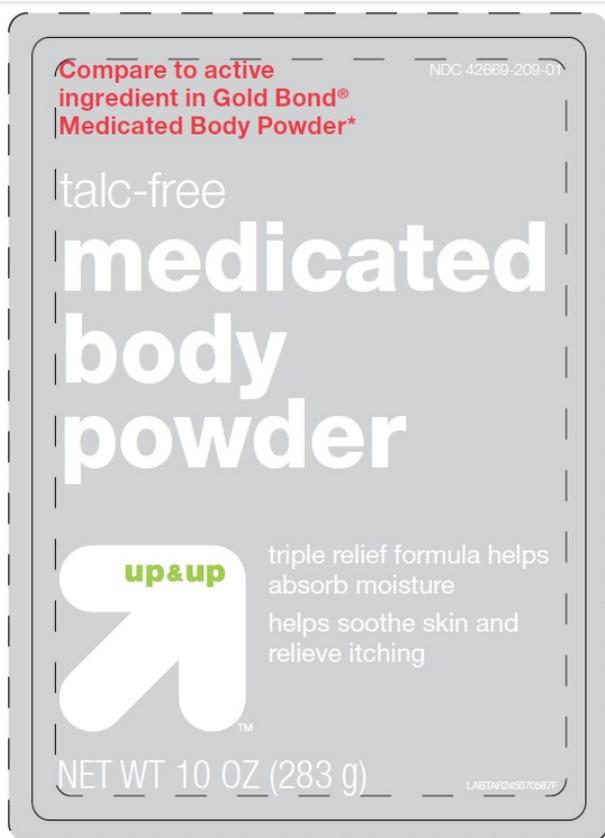
Compare to active ingredient in Gold Bond® Medicated Body Powder*

talc-free medicated body powder

triple relief formula helps absorb moisture

helps soothe skin and relieves itching

NET WT 10oz (283 g)



TALC-FREE MEDICATED BODY			
menthol powder			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42669-209
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)				
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
SALICYLIC ACID (UNII: O414PZ4LPZ)				
EUCALYPTUS OIL (UNII: 2R04ONI662)				
ZINC STEARATE (UNII: H92E6QA4FV)				
PEPPERMINT OIL (UNII: AV092KU4JH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42669-209-01	283 g in 1 BOTTLE; Type 0: Not a Combination Product	01/30/2018	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017		01/30/2018	

Labeler - Davion, Inc (174542928)

Revised: 12/2025

Davion, Inc