# TALC-FREE MEDICATED BODY- menthol powder Davion, Inc

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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 throughly 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

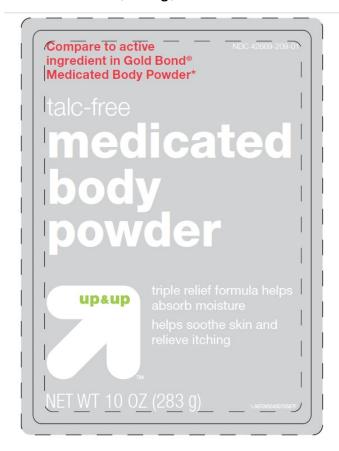
Compare to active ingredient in Gold Bond® Medicated Body Powder\*

talc-free medicated body powder

triple relief formulahelps 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
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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: 2R040Ni662)			
ZINC STEARATE (UNII: H92E6QA4FV)			
PEPPERMINT OIL (UNII: AV092KU4JH)			

ı	P	ackaging			
	#	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)

# Registrant - Davion, Inc (079536689)

Revised: 2/2024 Davion, Inc