

**HEB MEDICATED CORNSTARCH BODY POWDER- menthol powder  
Davion, Inc**

-----  
**H-E-B Medicated Cornstarch Body Powder Triple Relief Formula**

**Active Ingredient**

Menthol 0.15%

**Purpose**

External Analgesic

**Uses**

Temporary relief of pain and itch associate 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**

- 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, Tricalcium Phosphate, Sodium Bicarbonate, Zinc Oxide, Salicylic Acid, Zinc Stearate, Eucalyptus Oil, Peppermint Oil.

## **PRINCIPAL DISPLAY PANEL**

NDC 42669-217-10

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

H-E-B Medicated Cornstarch Body Powder

### **TRIPLE RELIEF FORMULA**

- Helps Absorbs Moisture
- Helps Soothe Skin
- Relieves Itching

Talc-Free

Net wt 10oz (283 g)

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

NDC #####



Medicated Cornstarch

# Body Powder

Triple Relief Formula

- Helps Absorb Moisture
- Helps Soothe Skin
- Relieves Itching

Talc-Free

NET WT. 10 OZ (283g)

LABHEB000000F

4.25"

## HEB MEDICATED CORNSTARCH BODY POWDER

menthol powder

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42669-217
Route of Administration	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.15 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TRICALCIUM PHOSPHATE</b> (UNII: K4C08XP666)	
<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z)	
<b>SALICYLIC ACID</b> (UNII: O414PZ4LPZ)	
<b>ZINC STEARATE</b> (UNII: H92E6QA4FV)	
<b>EUCALYPTUS OIL</b> (UNII: 2R04ONI662)	
<b>PEPPERMINT OIL</b> (UNII: AV092KU4JH)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42669-217-10	283 g in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2018	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/22/2018	

**Labeler** - Davion, Inc (174542928)**Registrant** - Davion, Inc (079536689)