H-E-B MEDICATED CORN STARCH FOOT POWDER- menthol powder
Davion, Inc

H-E-B Medicated Corn Starch Foot Powder

Active Ingredient

Menthol 1.0%

Purpose

External Analgesic

Uses

Temporary relief of pain and itch associate with:

- Minor Cuts
- Sunburn
- Insect Bites
- Scrapes
- Minor Burns
- Minor Skin Irritations

Warnings

- **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, Benzethonium Chloride, Eucalyptus Oil, Peppermint Oil.

PRINCIPAL DISPLAY PANEL

NDC 42669-218-10

Compare to Gold Bond ® Medicated Foot Powder active ingredient*

H-E-B Medicated Corn Starch Foot Powder

TRIPLE RELIEF FORMULA

- Controls Odor
- Absorbs Moisture
- Relieves Itching

Talc-Free

NET WT. 10 OZ (283g)

H-E-B MEDICATED CORN STARCH FOOT POWDER

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42669-218
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)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42669-218-10	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)

Revised: 2/2024 Davion, Inc