

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)

Compare to Gold Bond® Medicated Foot Powder active ingredient**

NDC #####



Medicated Cornstarch

Foot Powder

Triple Relief Formula

- Controls Odor
- Absorbs Moisture
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NET WT. 10 OZ (283g)

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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)

Revised: 12/2025

Davion, Inc