

**MAXIMUM STRENGTH MEDICATED FOOT POWDER- menthol powder  
Davion, Inc**

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**MAXIMUM STRENGTH Medicated Foot Powder with Corn Starch**

**Active Ingredient**

Menthol 1.0%

**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

- 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, Tricalcium Phosphate, Sodium Bicarbonate, Benzethonium Chloride, Eucalyptus Oil, Peppermint Oil

## PRINCIPAL DISPLAY PANEL

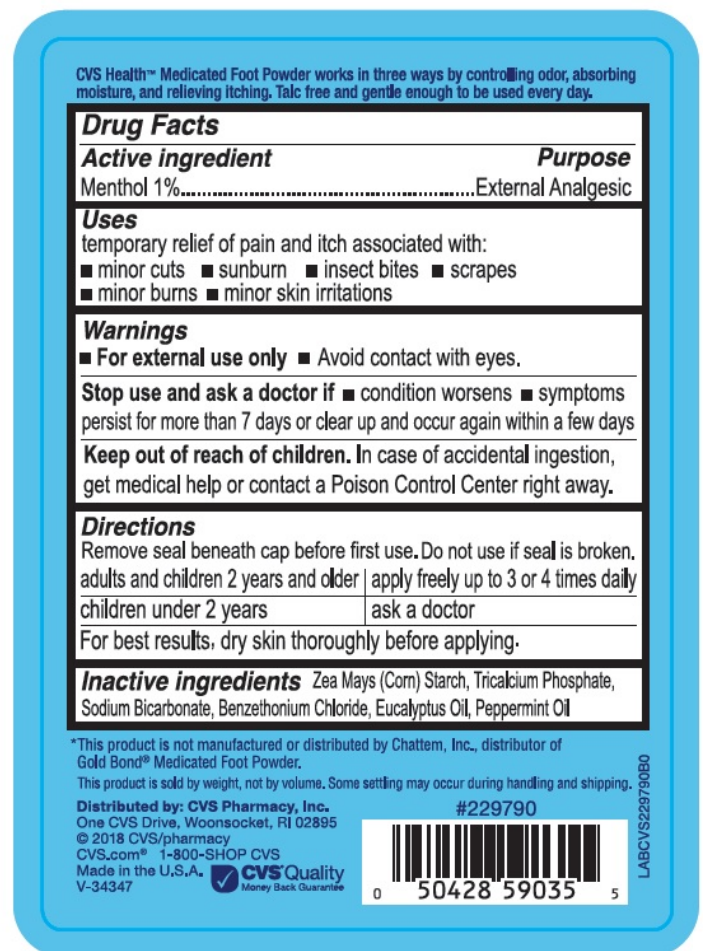
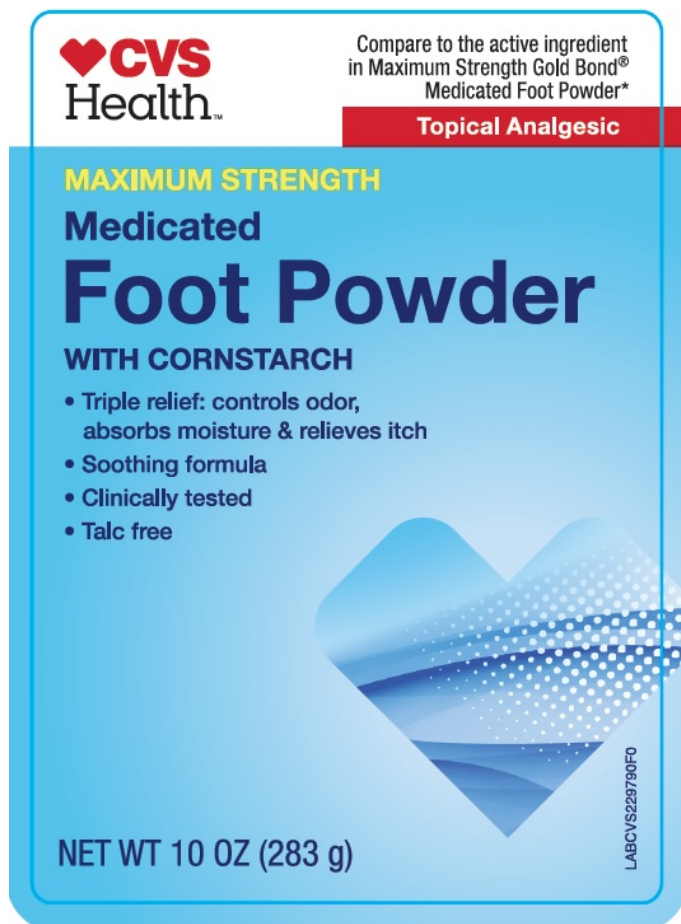
NDC 42669-283-01

CVS HEALTH Compared to the active ingredient in Maximum Strength Gold Bond Medicated Foot Powder\*

MAXIMUM STRENGTH Medicated Foot Powder WITH CORNSTARCH

- Triple Relief: controls odor, absorbs moisture & relieves itc
- Soothing formula
- Clinically tested
- Talc free

NET WT 10 OZ (283 g)



## MAXIMUM STRENGTH MEDICATED FOOT POWDER

menthol powder

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:42669-283
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**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)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42669-283-01	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