WALGREENS MEDICATED FOOT POWDER- menthol powder
Davion, Inc

Walgreens Foot Powder

Active Ingredient

Menthol 1.0%

Purpose

External Analgesic

Uses

For the temporary relief of pain and itch associate with minor skin irritations on the foot

Warning

- For external use only

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- condition worsens
- symptoms persists for more than 7 days or clear up and occur again within a 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

  Thoroughly wash and dry feet, sprinkle powder liberally over feet, between toes and on bottom of feet and in shoes

Inactive ingredients

Talc, Sodium Bicarbonate, Acacia, Benzethonium Chloride, Eucalyptus Oil, Peppermint Oil

Principal Display Panel

NDC 42669-100-10

WALGREENS FOOT POWDER

Compare to Gold Bond Medicated Foot Powder active ingredient

Foot Powder

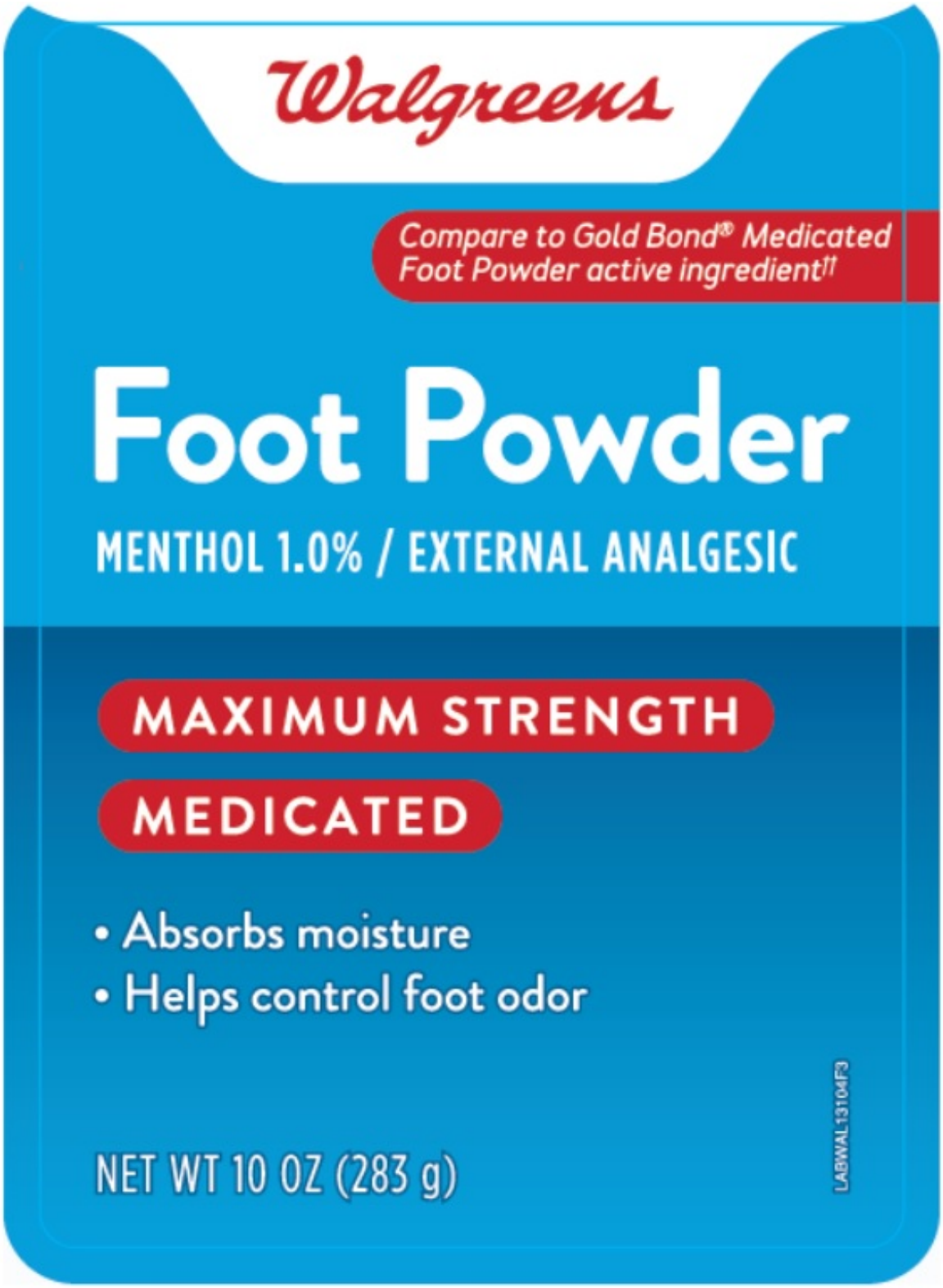
MENTHOL 1.0%/ EXTERNAL ANALGESIC

MAXIMUM STRENGTH

MEDICATED

Absorbs Moisture
Helps control foot odor

NET WT 10 OZ (283 g)



WALGREENS MEDICATED FOOT POWDER

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42669-100
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
PEPPERMINT OIL (UNII: AV092KU4JH)	
TALC (UNII: 7SEV7J4R1U)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
ACACIA (UNII: 5C5403N26O)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42669-100-10	283 g in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	06/01/2018	

Labeler - Davion, Inc (174542928)

Registrant - Davion, Inc (079536689)

Revised: 2/2024 Davion, Inc