MENTHOL- maximum strength medicated foot powder powder
Harmon Store Inc.

Harmon Medicated Foot Powder

Active ingredient

Menthol 1.0%

Purpose

External analgesic

Use

for the temporary relief of pain and itching associated with minor skin irritation on the foot

Warnings

For external use only.

When using this product
- avoid contact with eyes

Stop and consult a doctor if
- conditions 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 of age and older, apply to affected area not more than 3 to 4 times daily
- children under 2 years of age, consult a doctor
- wash and dry feet thoroughly
- sprinkle powder liberally on feet, between toes and on bottoms of feet

Inactive ingredients

benzethonium chloride, eucalytus oil, gum acacia, peppermint oil, sodium bicarbonate, talc

Questions?

Call 1-866-964-0939

Principal Display Panel

Harmon

FACEVALUES

MAXIMUM STRENGTH

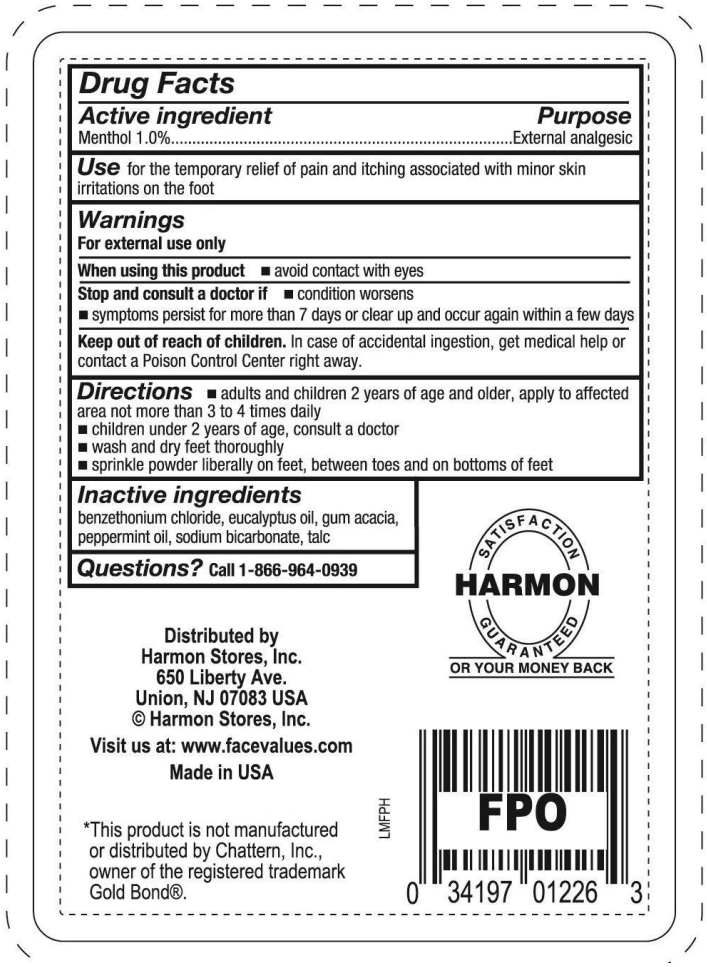
Medicated

Foot Powder

Menthol 1%

Triple Relief Formula
- Absorbs Moisture
- Relieves Itching
- Controls Foot Odor

NET WT 10 OZ (283 g)



MENTHOL

maximum strength medicated foot powder powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63940-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.8 g in 283 g

Inactive Ingredients

Ingredient Name	Strength
ACACIA (UNII: 5C5403N26O)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TALC (UNII: 7SEV7J4R1U)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63940-014-10	283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/18/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2012	

Labeler - Harmon Store Inc. (804085293)

Revised: 2/2024 Harmon Store Inc.