MAXIMUM STRENGTH MEDICATED FOOT POWDER- medicated foot powder powder
Walmart Stores Inc

Equate Medicated Foot Powder Menthol 1%

Active ingredient

Menthol 1.0%

Purpose

External analgesic

Use

for the temporary relief of pain and itching associated with

-minor cuts

-scrapes

-sunburn

-insect bites

-prickly heat

-rashes

-minor burns

-minor skin irritation

-dries the oozing of poison ivy, oak and sumac.

Warnings

For external use only.

When using this product

- avoid contact with eyes. Not for genital area

Stop and consult a doctor if

- conditions worsens
- redness, irritation, swelling or pain persist or increases
- symptoms do not get better within 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 (1-800-222-1222) right away.

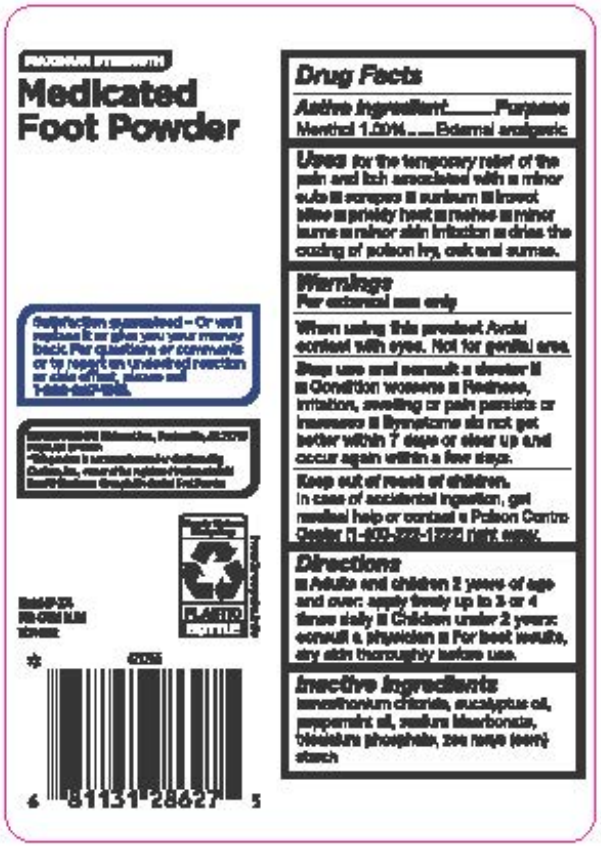
Directions

- adults and children 2 years of age and over: apply freely up to 3 or 4 times daily
- children under 2 years: consult a physician
- For best results, dry skin thoroughly before use.

Inactive ingredients

benzethonium chloride, eucalyptus oil, peppermint oil, sodium bicarbonate, tricalcium phosphate, Zea mays (corn) starch

Maximum Strength Medicated Foot Powder



MAXIMUM STRENGTH MEDICATED FOOT POWDER

medicated foot powder powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-795
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.1 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
ZEA MAYS WHOLE (UNII: 1G5HNE09V8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-795-01	283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	09/01/2020	

Labeler - Walmart Stores Inc (051957769)

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