#### MAXIMUM STRENGTH MEDICATED FOOT POWDER- medicated foot powder powder Target Corporation

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# Target 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 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 1-800-910-6874

# **Principal Display Panel**

Maximum strength

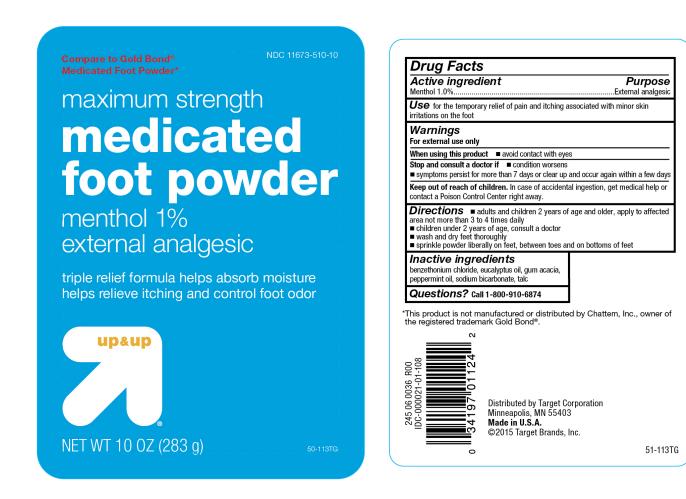
### medicated

#### foot powder

menthol 1% external analgesic

triple relief formula helps absorb moisture helps relieve itching and control foot odor

NET WT 10 OZ (283g)



TOPICAL   TOPICAL   Active Ingredient/Active Moiety   Ingredient Name Basis of Strength Strength   MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL 0.1 g in 1 g   Ingredient Name Strength   Ingredient Name Strength   BENZETHONIUM CHLORIDE (UNII: PH41D05744)   EUCALYPTUS OIL (UNII: SC5403N260)   FEPPERMINT OIL (UNII: AV092KU4JH)   Sociul BICARBONATE (UNII: 8MDF5V39QO)   TALC (UNII: 7SEV7J4R1U)   Packaging   # tem Code Package Description Marketing Start Marketing Enc   Date   ACC11673- 283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Od/30/2013	medicated foot	powder powd	er				
Notice of Administration TOPICAL   TOPICAL   Active Ingredient/Active Moiety   Ingredient Name Basis of Strength Strength   MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL 0.1 g in 1 g   Ingredient Name Basis of Strength   Ingredient Name Strength   BENZETHONIUM CHLORIDE (UNII: PH41D05744)   EUCALYPTUS OIL (UNII: 2R040NI662)   AcACIA (UNII: SC5403N260)   PEPPERMINT OIL (UNII: AV092KU4JH)   Social Marketing Start (UNII: 80DF5V39Q0)   TALC (UNII: 7EV7J4R1U)   Packaging   # Marketing Information   Marketing Information   Marketing Combination Product	Product Info	ormation					
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Ingredient Name   Basis of Strength   Strength     MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)   MENTHOL   0.1 g in 1 g     Inactive Ingredients   Ingredient Name   Strength     Ingredient Name   Strength   Strength     Ingredient Name   Strength   Ingredient Name     Ingredient Name   Strength   Ingredient Name     Ingredient Name   Strength   Strength     Ingredient Name   Strength   Strength     Ingredient Name   Strength   Ingredient Name     Ingredient Name   Strength   Ingredient Name     Strength   Strength   Ingredient Name   Ingredient Name     Stolun BicARBONATE (UNII: StrengtNamag)   Pa	Route of Admi	nistration	TOPICAL				
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Marketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End Date							
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OTC Monograph Drug   M017   04/30/2013		Applica		raph		Marketing End Date	
	OTC Monograph [	Drug M017		0	4/30/2013		

Labeler - Target Corporation (006961700)

Revised: 2/2024

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