MEDICATED FOOT POWDER- menthol powder Rejoice International Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Medicated Foot Powder

Active Ingredients Purpose Menthol 1.0% Anti-itch

∏Uses

temporarily relieves the pain and itch associated with minor skin irritation on the foot

- provides itch relief
- absorbs excess moisture
- helps control foot odor and odor-causing bacteria
- cools and soothes irritated skin

Keep out of reach of children. In case of accidental ingestion, get medical help or contact a Poison Control Center right away.

XtraCare

Medicated Foot Powder

3x Action

Itch Relieving

Odor Controlling

Moisture Absorbing

□Directions

adults and children 2 years and older

apply freely up to 3 or 4 times daily

children under 2 years

ask a doctor before using

Thoroughly wash and dry feet, sprinkle powder liberally over feet, between toes, and on bottom of feet. For best results apply to affected areas up to 3 or 4 times daily. Sprinkle inside shoes for maximum freshness.

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□Warnings

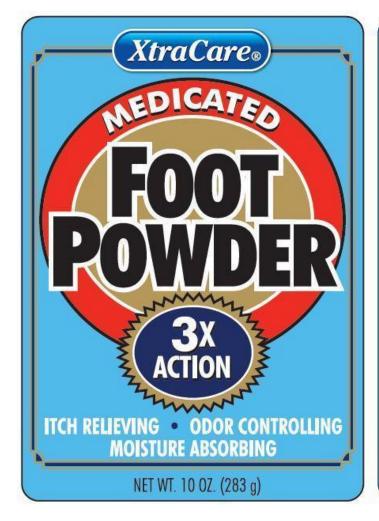
For external use only

• avoid contact with eyes

Stop using and ask a doctor if

- condition worsens
- symptoms do not get better within 7 days

☐ *Inactive ingredients* talc, sodium bicarbonate, acacia, benzethonium chloride, eucalyptus oil, peppermint oil





MEDICATED FOOT POWDER

menthol powder

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:57337-078

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
	TALC (UNIII 705) (714D1U)	

TALC (UNII: 7SEV7J4R1U)

SODIUM BICARBONATE (UNII: 8MDF5V39QO)

ACACIA (UNII: 5C5403N260)

BENZETHONIUM CHLORIDE (UNII: PH41D05744)

EUCALYPTUS OIL (UNII: 2R040NI662)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

l	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
l		NDC:57337-078- 01	283 g in 1 PACKAGE; Type 0: Not a Combination Product	06/15/2015			

Marketing Information						
Marketing Category			Marketing End Date			
OTC monograph not final	part348	06/15/2015				

Labeler - Rejoice International Inc. (078741245)

Registrant - Jell Pharmaceuticals Pvt Ltd. (726025211)

Establishment						
Name	Address	ID/FEI	Business Operations			
Jell Pharmaceuticals Pvt. Ltd.		726025211	manufacture(57337-078)			

Revised: 11/2022 Rejoice International Inc.