

MAXIMUM STRENGTH MEDICATED FOOT POWDER- medicated foot powder powder
Jell Pharmaceuticals Pvt. Ltd.

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.

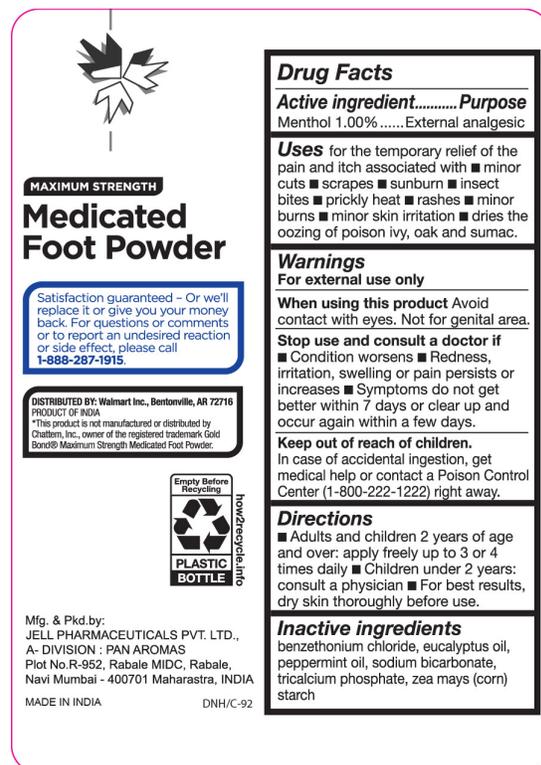
Inactive ingredients

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

Maximum Strength Medicated Foot Powder

MAX STRENGTH FOOT POWDER 10oz

SIZE : 76 x 108mm



Drug Facts

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medicated foot powder powder

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30400-507	
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:30400-507-01	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	M016	09/01/2020		

Labeler - Jell Pharmaceuticals Pvt. Ltd. (726025211)

Registrant - Jell Pharmaceuticals Pvt. Ltd. (726025211)

Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharmaceuticals Pvt Ltd.		726025211	manufacture(30400-507)

Revised: 12/2024

Jell Pharmaceuticals Pvt. Ltd.