MEDICATED BODY- menthol and zinc oxide powder
Universal Distribution Center LLC

Medicated Body Powder

Drug Facts

Active Ingredient

Menthol 0.15%

Zinc Oxide 1.0%

Purpose

External analgesic

Skin Protectant

Uses

for the temporary relief of pain and itching associated with

• Backache • Minor Burn • Minor Skin Irritations

• Minor Cut • Sunburn • Insect Bites

• Ooozing and weeping of poison ivy, poison oak and poison sumac can be dried using Universal Medicated Powder

WARNING

for external use only.

When using this product avoid contact with the eyes.

Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- Adults and children 2 years of age and older: apply to affected area no more than 3 to 4 times daily
- Children under 2 years of age: consult a doctor
- For best results, dry area thoroughly before applying

Inactive Ingredients

Zea mays (corn) starch, sodium bicarbonate, tricalcium phosphate, acacia seyal gum, eucalyptol, methyl salicylate, salicylic acid, thymol, zinc stearate

WITH ESSENTIAL OILS

Daily Use Leaves Skin Feeling Fresh & Healthy

PURE CORNSTARCH

This product is sold by weight, not by volume. Some settling may occur during handling and shipping.

Distributed By:
Universal Distribution Center
Edison, NJ 08817
www.universaldc.com

Made in India

Packaging

MEDICATED BODY
menthol and zinc oxide powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-038
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	1.5 mg in 1 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
GUM TALHA (UNII: H18F76G097)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
THYMOL (UNII: 3J50XA376E)	
ZINC STEARATE (UNII: H92E6QA4FV)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-038-25	283 g in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2017	
2	NDC:52000-038-26	226 g in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/04/2017	

Labeler - Universal Distribution Center LLC (019180459)

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