

BODY- menthol powder

Rite Aid

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.

Drug Facts

Active ingredient

Menthol 0.15%

Purpose

External analgesic

Uses

for temporary relief of pain and itching due to:

- minor cuts
- sunburn
- insect bites
- poison ivy
- poison oak
- poison sumac
- scrapes
- minor burns
- minor skin irritations

Warnings

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 right away.

Directions

- adults and children 2 year of age and older - apply to affected area not more than 3 to 4 times daily
- children under 2 years of age - do not use, consult a doctor
- for best results dry skin thoroughly before applying

Inactive ingredients

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

disclaimers

This product is not manufactured or distributed by Chattem, distributor of Gold Bond Medicated Body Powder.

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

Adverse Reaction

DISTRIBUTED BY: RITE AID

30 HUNTER LANE

CAMP HILL, PA 17011

100% GUARANTEE

or your money back.

379.001/379AB

Principal Display Panel

Compare to active ingredient of Gold Bond Medicated Body Powder

RITE

AID

RENEWAL

BODY POWDER

External Analgesic

MEDICATED

Itch Relieving

Cooling, Absorbent

NET WT 10 OZ (283 g)

*Compare to active ingredient of Gold Bond Medicated Body Powder



RENEWALTM

BODY POWDER
External Analgesic

MEDICATED

Itch Relieving,
Cooling, Absorbent



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L0013927FA

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RENEWALTM

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BODY

menthol powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0000
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength

MENTHOL (UNII: L7T10EP3A) (MENTHOL - UNII:L7T10EP3A)			MENTHOL	1.5 mg in 1 g
Inactive Ingredients				
Ingredient Name			Strength	
STARCH, CORN (UNII: O8232NY3SJ)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)				
ZINC OXIDE (UNII: SOI2LOH54Z)				
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:11822-0000-3	283 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/19/2013	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part348	11/19/2013	

Labeler - Rite Aid (014578892)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
MK Packaging		047022405	manufacture(11822-0000)

Revised: 4/2018

Rite Aid