BODY- menthol, zinc oxide powder
Target Corporation

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.

UP & UP 379.001-379AB

Active ingredient

Menthol 0.15%

Purpose

External analgesic

Uses

for temporary relief of pan and itching due to:

- mnor 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 symtpoms 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

talc, zinc oxide, acacia seyal gum, eucalyptol, methyl salicylate, salicylic acid, thymol, zinc stearate

Questions

Call 1-800-910-6874

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 Target Corporation

Minneapolis, MN 55403

Made in the U.S.A. with U..S and foreign components

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Compare to Gold Bond

Medicated Body Poweder

medicated

body

powder

external analgesic

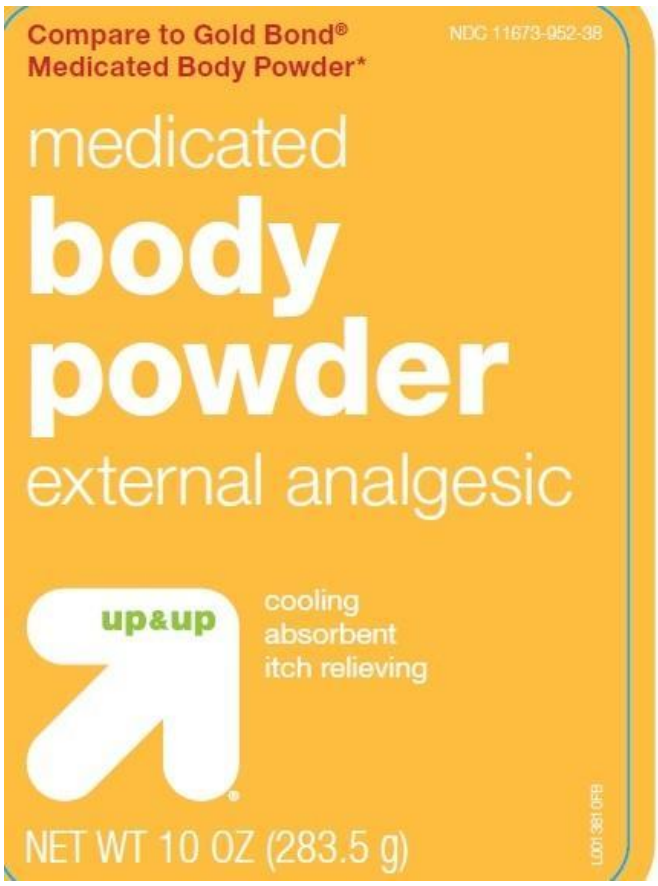
up & up

cooling

absorbant

itch relieving

NET WT 10 OZ (283.5 g)



BODY

menthol, zinc oxide powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-952
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.5 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
TALC (UNII: 7SEV7J4R1U)	
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:11673-952-38	283 g in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/01/2013	

Labeler - Target Corporation (006961700)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(11673-952)

Revised: 4/2022 Target Corporation