BODY- menthol powder
Supervalu

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 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 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 years 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

Questions

Call 1-877-932-7948

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 SUPERVALU INC

EDEN PRAIRIE, MN 55344 USA

We're committed to your satisfaction and guarantee the quality of this product.

Contact us at 1-877-932-7948 or www.supervalu-ourownbrands.com

Please have package available.

379.001/379AB

Principal Display Panel

EQUALINE

Compare to Gold Bond Medicated Body Powder

MEDICATED

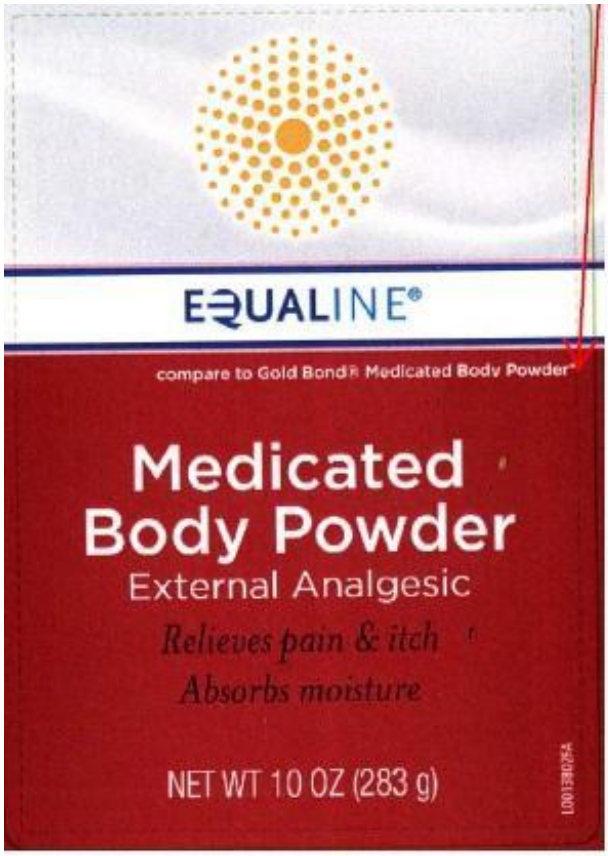
BODY POWDER

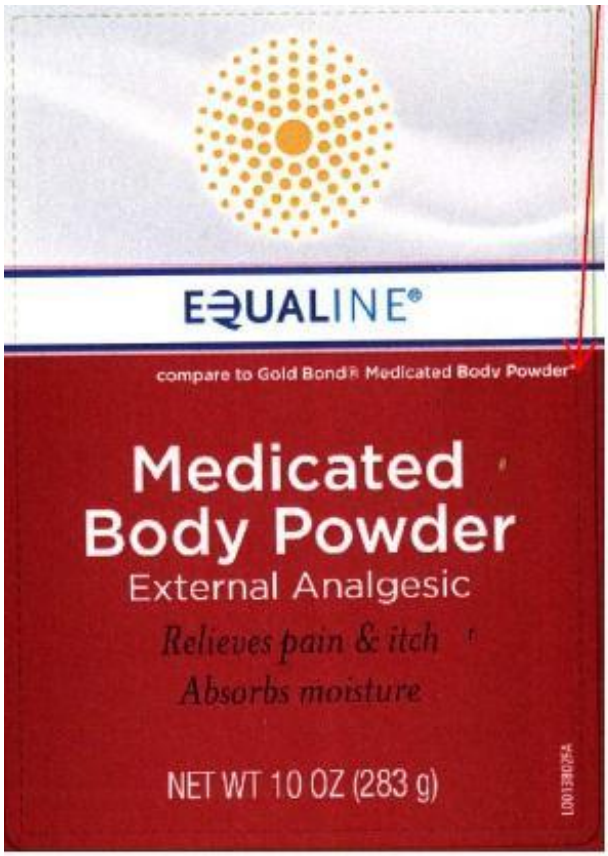
External Analgesic

Relieves pain & itch

Absorbs moisture

NET WT 10 oz (283g)





BODY
menthol powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-003
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
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:41163-003-38	283 g in 1 BOTTLE; Type 0: Not a Combination Product	09/27/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/27/2013	

Labeler - Supervalu (006961411)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
MK Packaging		047022405	manufacture(41163-003)

Revised: 3/2018 Supervalu